Title of the Invention

INHIBITING THE GROWTH OF BACTERIAL BIOFILMS

Cross-Reference to Related Applications

The present invention claims the benefit of U.S. provisional application no.

5   60/464,333 filed April 22, 2003 and U.S. provisional application no. 60/517,391 filed

November 6, 2003, the entire contents of both applications are incorporated herein by

reference.

BACKGROUND OF THE INVENTION

Field of the Invention

10   The present invention relates to the inhibition or reduction in bacterial biofilm

growth and development.

Discussion of the Background

Surface attached, matrix-enclosed communities, called biofilms, cause serious

economic and health problems due to biofilm-associated phenotypes such as antibiotic

15   resistance or biofouling Costerton, J. W., Stewart, P. S. & Greenberg, E. P. (1999)

Science 284, 1318-22. The inherent resistance to antimicrobial agents are the root of

many persistent and chronic bacterial infections as nosocomial infections and legionaire's

disease. The drastic phenotypic changes seen in biofilms led to the assumption that the

physiological modifications necessary for planktonic bacteria to adopt the biofilm

20   lifestyle must involve specific responses. However, biofilm physiology is still poorly

understood and, whereas the early events of biofilm formation are well documented, little

is known about the nature of the physiological changes and critical regulatory processes

occurring inside mature biofilms. Global expression profiling comparing protein

synthesis in *Pseudomonas* planktonic and biofilm bacteria suggested that a large number

25   of genes could be differentially regulated during biofilm development (Sauer, K., Camper,

A. K., Ehrlich, G. D., Costerton, J. W. & Davies, D. G. (2002) *J Bacteriol* **184,** 1140-54; Sauer, K. & Camper, A. K. (2001) *J Bacteriol* **183,** 6579-89; Whiteley, M., Bangera, M. G., Bumgarner, R. E., Parsek, M. R., Teitzel, G. M., Lory, S. & Greenberg, E. P. (2001) *Nature* **413,** 860-4). Although these pioneering studies opened the way to the genetic

5   characterization of the biofilm phenotype, extracting functional information from genomic approaches remains a challenge.

Escherichia coli K12, a widely used bacterial model, does not spontaneously form extensive biofilms. However, it has been previously shown that expression of pili from conjugative plasmids, which are widespread in natural bacterial populations, promotes

10   the development of mature biofilms (Ghigo, J. M. (2001) *Nature* **412,** 442-5). This raised the possibility of studying the genetic basis of the biofilm phenotype in *E. coli* K12 where expression profiling can be combined with the phenotypic analysis of a large set of deletion mutants.

In view of the above, there remains an urgent need to develop new strategies for

15   combating the development of mature biofilms Based on the discovery of the genes involved in the development of mature biofilms, the present invention provides targets to disrupt the development, formation and/or maturation of bacterial biofilms, and molecular tools to characterize and detect mature biofilms.

20   SUMMARY OF THE INVENTION

Thus, the present invention is based on the discovery of the unique expression of genes during the formation of bacterial biofilms thereby providing a target to reduce, ameliorate, attenuate, inhibit and/or treat biofilms.

Accordingly, one aspect of the present invention is to a method of treating,

25   reducing, ameliorating, attenuating and/or inhibiting the formation of biofilms by

targeting the specific genes that are involved in the formation of the biofilm. These methods can be accomplished by contacting an already formed biofilm and/or a sample, surface or other substrate that may be susceptible to biofilm formation with one or more inhibitors of those genes.

5   In another aspect of the present invention, methods of screening for substances that inhibit the genes involved in biofilm formation is also provided.

In another aspect of the present invention, a polynucleotide library which is useful for molecular characterization of a mature bacterial biofilms is also provided.

In another aspect of the present invention, using the libraries to detect mature

10   bacterial biofilms is also provided.

BRIEF DESCRIPTION OF THE DRAWINGS

A more complete appreciation of the invention and many of the attendant advantages thereof will be readily obtained as the same becomes better understood by reference to the following detailed description when considered in connection with the

15   accompanying drawings, wherein:

Figure 1: Function of genes over-expressed in TG1 biofilm versus exponential growth phase

This figure summarizes the data presented in Table 3. The genes have been classified according to the COGs functional categories annotation system. Large and medium size

20   numbers indicate the total number of *E. coli* biofilm-induced genes into each class or sub-class of indicated functions. Genes are indicated only when their expression level in biofilm differed by at least a two-fold factor (≥ 2). Numbers within brackets indicate the rank as over-expressed genes; 1 = most expressed gene in TG1 *E. coli* biofilm.

25   **Figure 2: Correlation of macroarray and quantitative real-time PCR results**

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The calculated macroarray and Q-RT-PCR ratios of the expression of 7 genes in TG1 biofilm relative to exponential growth phase were log transformed, and values were plotted against each other to evaluate their correlation. The correlation coefficient was deduced from a linear regression of the plotted values.

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Figure 3: Biofilm phenotype of selected deletion mutants

Mature biofilm development of *E. coli* TG1 (wt) compared with a selection of deletion mutants of genes over-expressed in TG1 biofilm.

A: For each mutant phenotype analysis, the extent of biofilm formation is shown in the

10   bottom part of the micro-fermenter and on the removable glass slide. A typical experiment is shown.

B: Graphical comparison of biofilm formation relative to wild type from the mutants presented in A. Data represents the average of three independent experiments for each mutant. The level of biofilm formed by wt TG1 biofilm was set to 100%.

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Figure 4: Functional profiling of *E. coli* biofilm: flow chamber analysis

A. Spatial distribution of biofilm formation for *E. coli* TG1 and selected TG1 deletion mutants expressing Gfp. Biofilms were grown in flow chambers. Biofilm development

20   was monitored by SCLM at the indicated times after inoculation (20 h, 45 h, 70 h, 95 h). Micrographs represent simulated three-dimensional images. Images inseted into 70 h and 95 h of *ycfJ* correspond to rare area where the biofilm was more developed.

B. COMSTAT analysis of biofilm structures. Diagrams and standard deviations (numbers indicated in the individual columns) of biomass and substrate coverage from biofilms of

25   *E. coli* TG1 and TG1 deletion mutants were determined by the COMSTAT program at

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four different time points (20 h, 45 h, 70 h, 95 h). Values are means of data from 12 image stacks (6 image stacks from two independent channels). The biomass is in the unit $\mu m^3/\mu m^2$. The substratum coverage values are relative (1 represents total coverage).

Figure 5: A comparison of biofilm formation capacity of mutants in the *E. coli cpx* and *rpoE* envelope stress pathways

Biofilm development comparison of TG1 and TG1 deletion mutants in micro-fermenters. The average of at least four experiments was plotted in the histogram. The level of biofilm formed by wt TG1 biofilm was set to 100%.

Figure 6: Comparison of TG1 and TG1 Δ*cpxP* biofilm structure

Phenotypic analysis of the structure of TG1 and TG1Δ*cpxP* biofilms grown in micro-fermenter.

A: General view of the bottom part of the fermenter.

B: Macroscopic biofilm grown on the internal glass slide, removed from the fermenter shown in panel A.

C: Close-up on the biofilm shown in panel B.

D: transverse section of TG1 and TG1Δ *cpxP* biofilm.

E and F: detailed X50 and X 10000 electron micrographs of TG1 and TG1 *cpxP* biofilm structure.

Figure 7: COG functional classes for genes under-expressed in TG1 biofilm versus exponential growth phase.

This figure summarizes the data presented in Table 4. The genes have been classified according to the COGs functional categories annotation system. Large and medium size numbers indicate the total number of *E. coli* genes falling into each class or sub-class of

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function. Genes are indicated only when their expression level in biofilm differed by at least a two-fold factor (≤ 0.5). Numbers within brackets indicate the rank as under-expressed genes; 1 = most repressed gene in TG1 biofilm.

5 **Figure 8: Functional profiling of mature *E. coli* biofilm: biofilm formation in microfermenters.**

Comparison of mature biofilm development in micro-fermenters of wild type *E. coli* TG1 with TG1 mutants in the genes found to be induced by over a two-fold factor in TG1 biofilm. This figure complements the Figure 3. The far right of the panel describes the

10 analysis of biofilm development of a *pspF* mutant, a constitutively expressed positive regulator of the *pspABCDE* operon. The data represent the average of at least three independent experiments for each mutant. Wild type TG1 biofilm formation was set to 100.

15 **Figure 9: Functional profiling of early steps in *E. coli* biofilm formation**

Comparison of the early adhesion ability of TG1 mutants in genes identified as over-expressed in mature TG1 versus exponential growth phase or analyzed in this study as visualized by crystal violet staining in a static microtiter plate-based assay. *E. coli* TG (M63B1 glucose medium supplemented with proline) adheres poorly in this assay. TG1

20 *fimA* (boxed) displays an expected reduced early adhesion capacity. Stars (*) correspond to TG1 mutants with a growth impairment leading to a non meaningful reduction of adhesion in this early biofilm assay.

25 DETAILED DESCRIPTION OF THE INVENTION

The formation of biofilms results in a major lifestyle switch that is thought to affect the expression of multiple genes and operons. Using DNA arrays to study the global effect of biofilm formation on gene expression, the inventors have demonstrated that in biofilms, 1.9% of the genes showed a consistent up or down-regulation by a factor greater than two, and that 10% of the E. Coli genome is significantly differentially expressed including genes of unknown function, stress-response genes as well as energy production and envelope biogenesis functions. The inventors provide evidence that the expression of stress envelope response genes, such as the *psp* operonor elements of the *cpx* pathway, is a general feature of *E. coli* biofilms. Using gene disruption of 53 of the genes showed that 17 of the genes are required for the formation of mature biofilm. This includes 11 genes of previously unknown function.

Thus, the genes involved in biofilm formation and useful as targets for identifying substances that inhibit biofilm formation are those described herein, for example, including *lctR, recA, mdh, rbsB, msrA, finA, tatE, pspF, cpxP, spy, ycfJ, ycfR, yoaB, yqcC, yggN, ymcA, yccA, yfcx, yghO, yceP,* and *ycuB.* Preferably, the genes involved in biofilms formation are one or more of yccA, (SwissProt accession number-P06967; GenBank number- g1787205, the amino acid sequence is shown as SEQ ID NO:299 and the nucleotide sequence encoding the protein is shown in SEQ ID NO:300), ycfJ, (SwissProt accession number- P37796;GenBank number-g1787353, the amino acid sequence is shown as SEQ ID NO:301 and the nucleotide sequence encoding the protein is shown in SEQ ID NO:302), and yceP, (SwissProt accession number-P75927;GenBank number-g1787299, the amino acid sequence is shown as SEQ ID NO:303 and the nucleotide sequence encoding the protein is shown in SEQ ID NO:304).

As used herein, the term "polynucleotide" refers to a polymer of RNA or DNA that is single-stranded, optionally containing synthetic, non-natural or altered nucleotide

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bases. A polynucleotide in the form of a polymer of DNA may be comprises of one or more segments of cDNA, genomic DNA or synthetic DNA.

The term "subsequence" refers to a sequence of nucleic acids that comprise a part of a longer sequence of nucleic acids.

5   In a further embodiment of the invention, the proteins are at least 70%, preferably at least 80%, more preferably at least 90% identical to the sequences identified above. In another embodiment, the genes and thus gene products that are to be inhibited are encoded by polynucleotide sequence with at least 70%, preferably 80%, more preferably at least 90%, 95%, and 97% identity to the sequences described above, these

10   polynucleotides will hybridize under stringent conditions to the coding or non-coding polynucleotide sequence above. Preferably, these homologous sequences would have the same or similar activity to the sequences specifically identified above.

The terms "stringent conditions" or "stringent hybridization conditions" includes reference to conditions under which a polynucleotide will hybridize to its target sequence,

15   to a detectably greater degree than other sequences (e.g., at least 2-fold over background). Stringent conditions will be those where hybridization in 50% formamide, 1 M NaCl, 1% SDS at 37°C., and a wash in 0.1X SSC at 60 to 65°C (see Tijssen, Laboratory Techniques in Biochemistry and Molecular Biology--Hybridization with Nucleic Acid Probes, Part I, Chapter 2 "Overview of principles of hybridization and the strategy of nucleic acid probe

20   assays", Elsevier, New York (1993); and Current Protocols in Molecular Biology, Chapter 2, Ausubel, et al., Eds., Greene Publishing and Wiley-Interscience, New York (1995)). Amino acid and polynucleotide identity, homology and/or similarity can be determined using the BLAST algorithm. Preferably, these homologous sequences would have the same or similar activity to the sequences specifically identified above.

In one embodiment, the present invention provides methods of reducing, inhibiting, ameliorating, and/or treating bacterial biofilms, such as *E. coli* biofilms, by inhibiting, reducing, and/or attenuating the genes and/or gene products, e.g, messenger RNA and proteins encoded thereby, described herein as being involved in biofilm

5   formation.

By "treating" is meant the slowing, interrupting, arresting or stopping of the progression of the biofilm growth and does not necessarily require the complete elimination of the biofilm. "Preventing" or "ameliorating" is intended to include the prophylaxis of the biofilm development and/or growth, wherein "prophylaxis" is

10   understood to be any degree of inhibition on the biofilm development and/or growth, including, but not limited to, the complete prevention of biofilm development and/or growth. The substances which inhibit the gene(s) described herein are collectively termed "biofilm inhibitor(s)." In one embodiment, the biofilm inhibitor(s) decrease the ability of the biofilm to develop and/or mature at least by 1%. In another embodiment,

15   the decrease is at least by 5%, 10%, 15%, 20%, 30%, 35%, 40%, etc.

To effectuate the inhibition of biofilms, a surface, and/or sample (collectively termed "at least one substrate") on which a biofilm has begun to develop can be contacted with one or more of the biofilm inhibitors thereby inhibiting the biofilm formation. In an alternative embodiment, the at least one substrate on which a biofilm has already formed

20   or developed can be contacted with one or more of the biofilm inhibitors such that biofilm becomes less prevalent or completely disappears from the substrate. In an alternative embodiment, the at least one substrate in which a biofilm has not begun to develop but is susceptible to biofilm formation can be pretreated with one or more of the biofilm inhibitors to inhibit the formation of the biofilm on the at least one substrate. The

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substrate as used herein refers to any surface, liquid or solid, on which a biofilm develops, has developed, or is susceptible to biofilm formation.

The biofilm inhibitors can be any substance, chemical, and/or biological materials that inhibit the development and/or formation of the biofilm in an appreciable manner as described herein. For example, antibodies that specifically bind to and inhibit the activity of proteins that are encoded by the genes described herein can be used to inhibit the development and/or formation of the biofilm. Polyclonal, monoclonal and/or fragments (e.g., Fab fragments) of antibodies that specifically bind to the proteins of the genes described herein may be used so long as they inhibit the function of the gene products according to the disclosure herein. Obtaining polyclonal, monoclonal and/or functional fragments thereof is conventional and is described, for example, in Harlow and Lane "Using Antibodies: A Laboratory Manual" © Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1999).

An effective amount of the inhibitors as described herein can be used either singularly or in combination and should be used in an amount that results in some inhibition of biofilm development and/or growth When the inhibitors are administered in combination, they may be premixed, administered simultaneously, or administered singly in series.

In another aspect of the present invention, methods to identify substances, agents, compounds and/or chemicals (collectively termed "inhibitors") that reduce, inhibit, ameliorate and/or treat the development of and/or formation of biofilms. Such methods are preferably accomplished by targeting one or more of the genes involved in biofilm development as described herein. In one embodiment of this aspect, the gene (or genes) are expressed in host cell, preferably a bacterial cell such as E. coli, and the ability of the inhibitor to affect the gene and/or protein are assessed. For example, levels of

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transcription can be measured using conventional DNA and/or RNA probing techniques, such as PCR and other hybridization assays. Thus, the cell expressing the one or more protein encoded by the genes described herein is contacted with the inhibitor and the relative level of transcription is measured in relation to the cell before contacting with the

5   inhibitor; and/or compared to a cell which similarly expresses the protein(s) and which was not contacted with the inhibitor. In a similar manner, levels of protein expressed in cell can be assessed, comparing contacted and uncontacted cells, using protein analytical techniques known in the art.

Screening for the inhibitors can also be accomplished by testing the effects of the

10   inhibitor(s) on the development and/or growth of the biofilm as described herein. Once identified, the inhibitors can be used to reduce, inhibit, ameliorate and/or treat the development of and/or formation of biofilms as described herein.

The inhibitors may be formulated or combined with any acceptable carrier, such as buffered saline or other buffered solution.

15   In another aspect of the invention, a polynucleotide library is provided that is useful in the molecular characterization of a mature bacterial biofilm, which comprises a pool of polynucleotide sequences or subsequences thereof wherein said sequences or subsequences are overexpressed in mature bacterial biofilms. The polynucleotide sequences or subsequences may be immobilized on a solid support in order to form a

20   polynucleotide array. As used herein, the term "immobilized on a support" means bound directly or indirectly thereto including attachment by covalent binding, hydrogen bonding, ionic interaction, hydrophobic interaction or otherwise. The solid support can be a nylon membrane, glass slide, glass beads, and/or a silicon chip. Thus, in another embodiment, a

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polynucleotide array is provided which is useful to detect a mature bacterial biofilm and which comprises an immobilized polynucleotide library as described above.

The immobilized polynucleotide library and array can be used for detecting differentially expressed polynucleotide sequences which are specifically correlated with a mature bacterial biofilms. In this method a polynucleotide sample is obtained, and labeled by reacting the polynucleotide sample with a labeled probe immobilized on a solid support wherein said probe comprises any of the polynucleotide sequences of the polynucleotide library as described above or an expression product encoded by any of the polynucleotide sequences; and detecting a polynucleotide sample reaction product. The method can be used for detecting mature bacterial biofilms, such as, an *Escherichia coli* biofilm.

In another embodiment of the method, a control polynucleotide sample, which is labeled, is employed for comparing the amount of polynucleotide sample reaction product to the amount of the control sample reaction product. In another embodiment of the method, RNA or mRNA is isolated from the polynucleotide sample, and which may be reverse transcribed to yield a cDNA molecule.

The labeling reaction can be performed by hybridizing the polynucleotide sample with the labeled probe. The label can be radioactive, colorimetric, enzymatic, molecular amplification, bioluminescent or fluorescent. Detection can then be performed as known in the art.

In another embodiment, where the product encoded by any of the polynucleotide sequences or subsequences is employed, the detection can be based on a receptor-ligand reaction.

In another aspect of the present invention, a method of detecting significantly overexpressed genes correlated with a mature bacterial biofilms can be performed. As used herein, "significantly overexpressed" means that the gene or expression product

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detected is expressed in by a factor of 2 or greater compared to a bacterial cell which is not in a biofilm or begun to develop biofilms characteristics. This method comprises detecting at least one polynucleotide sequence or subsequence of a polynucleotide library as described above or detecting at least one product encoded by said polynucleotide

5   library in a sample obtained from a patient. In another embodiment of this method, an amount of the at least one polynucleotide sequence or subsequence or product encoded by said polynucleotide sequence is compared with an amount of the polynucleotide sequence or subsequence or product encoded by said polynucleotide sequence or subsequence obtained from a control sample. Extracted mRNA may also be used, which can be

10   reverse transcribed into a cDNA molecule. In another embodiment of this method, the at least one polynucleotide sequence or subsequence can be hybridized with mRNA or cDNA from the polynucleotide sample using, for example, the labeling and detection described above. In another embodiment, of this method where the product encoded by any of the polynucleotide sequences or subsequences is employed, the detection can be

15   based on a receptor-ligand reaction.

Preferably, the sequences or subsequences correspond substantially to the polynucleotide sequences of the following genes : *rne, lctR, dinI, glpQ, mdh, sixA, lamB, rbsB, gadA, pspA, pspB, pspC, pspD, tatE, cpxP, rseA, rpoE, spy, yebE, yqcC, yfcX, yjbO, yceP,* and *ygiB*. In another embodiment, the library further comprises polynucleotide

20   sequences or subsequences thereof of the following genes: *recA, msrA, fimA, pspF, ycfJ, ycfR, yoaB, yggN, yneA, yccA, yghO*. In another embodiment, the library further comprises polynucleotide sequences or subsequences thereof of the following genes: *RplY, recA, cyoD, sucA, fdhF, cyoC, nifU, sucD, sfsA, nifS, fadB, ucpA, ftsL, sulA, eco, msrA, pspD, fimA, fimI, pspE, pspF, cutC, sodC, rseB, ycfJ, ycfR, yoaB, yhhY, yggN, yneA,*

25   *ybeD, ydcI, yddL, yccA, yrdD, ybjF, yihN, 1228, ycfL, yiaH, yqeC.*

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In another embodiment, the library further comprises polynucleotide sequences or subsequences thereof of the following genes: *lysU, miaA, rluC, rplY, crl, cspD, dniR, fruR, idnR, lacI, nac, rnk, rpoS, ttk, b0299, dinG, dinP, exo, intA, recA, recN, sbmC, xthA, aceA, aceB, aldA, atpA, cyoA, cyoC, cyoD, dctA, fdhF, fdoG, glpD, glpK, nifU, pckA,*

5   *sdhB, sdhD, sucA, sucB, sucD, xdhD, agp, gcd, glgS, glpX, malE, malF, malS, mglA, mglB, mrsA, pgm, rbsC, rbsD, sfsA, ansB, argC, argR, idnD, leuD, metH, nifS, putP, metK, pnuC, ubiE, fabA, fadB, fadE, fadL, pgpA, pssA, uppS, idnO, ucpA, ftsL, sulA, dnaJ, dnaK, eco, fkpA, glnE, htpG, htpX, msrA, amiB, ddg, fhiA, fimA, fimI, htrL, lepB, mraW, nlpB, nlpC, ompC, ompG, pspE, pspF, chaA, chaC, cutC, cysP, cysU, fur, modA, modB,*

10   *modC, modE, sodC, trkH, rseB, ycfJ, ycfR, yoaB, yhhY, yggN, yneA, ybeD, ydcI, yddL, yccA, yrdD, ybjF, yihN, ycfT, yeeF, yfiE, yeeD, yliH, yfcM, ybiX, yfhF/nifA, ygfQ, ybhR, ybdH, yihR, ydcT, ygiS, ybaZ, ydaM, tfaR, yceL, yheT, yjdC, ybiW, ybiF, ynaI, yceE, yhdP, ygjE, csiE, yfdE, yeeE, yegQ, glcA, yfdW, yfeT, ygjK, ydeW, b1228, ycfL, yghO, yiaH, yqeC, ycfT, yhjJ, yceB, ybiX, ygiQ, yagV, yoeA, ybhQ, ybcI, ybbF, ybgI, yncH, yfbM, yjiM,*

15   *yjfO, ychN, ynaC, ymfE, yfcN, yrbC, yfdQ, yfeY, ygiM, yhgA, yhjQ, yfcF, yfcI, yjiD, yfbP, yphB, yfbN, ylbH, ybhM, yrbL, yjfY, ynfA, yajI, yedI, yafZ, yjjU, yfhH, yafN, yrbE, yfgC, yfjQ, ycaK, yfeS, b4250, ybgA, yeeA, ypfI, b2394, yegK, ybcJ, yhiN, ypfG, ydiY, yjjJ, ycaP, yfgJ.*

In a preferred embodiment of the library, the biofilms is a *Escherichia coli*

20   biofilms.

EXAMPLES

Experimental procedures

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Bacterial strains and culture conditions

Bacterial strains used in this work are described in Table 2. All experiments were performed in 0.4% glucose M63B1 minimal medium at 37°C except flow chamber experiments that were performed at 30°C in 0,02% glucose FAB minimal medium.

5   Proline was added at 400 µg/ml for TG growth.

Early adhesion and biofilm formation assay

Microtiter plate assays were performed as described in (O'Toole, G.A., and Kolter, R. (1998) *Mol Microbiol* **30**: 295-304). Biofilm development comparisons in aerated

10   micro-fermenters were conducted as described in (Ghigo, J.M. (2001) *Nature* **412**: 442-445.). The biofilms formed on the removable glass slide were photographed and then resuspended in 10 ml of M63B1 minimal medium. The optical density at 600 nm (OD_{600}) of the resuspension was then measured. After 24 hours the average resuspended *E. coli* TG1 biofilm biomass reached $OD_{600}= 5$. Each mutant was tested in at least 3 independent

15   experiments alongside with the control strain TG1.

Macroarray analysis

Genomic expression profiles were performed on *E. coli* TG1 and TG strains grown in 0.4% glucose M63B1 at 37°C either as planktonic cultures or mature biofilms. Planktonic

20   cultures were realized in agitated Erlenmeyer flasks (main experiment) or aerated micro-fermenters, both in exponential phase $OD_{600} \sim 0.6$ or stationary phase $OD_{600} \sim 3$. Mature biofilms were grown in aerated micro-fermenters (8 and 5 day old biofilms for TG1 and TG respectively). For all conditions, the equivalent of 15 OD_{600} of bacterial cells was collected. The cells were then broken in a Fast Prep apparatus (Bio 101). Total RNA

25   was extracted by Trizol (Gibco-BRL) treatment. Genomic DNA was degraded using the

DNA-*free*™ kit (Ambion). Radioactively labeled cDNAs, generated by using *E. coli* K12 CDS-specific primers (SIGMA-GenoSys), were hybridized to *E. coli* K12 panorama gene arrays containing duplicated spots for each of the 4,290 predicted *E. coli* K12 ORFs (SIGMA-GenoSys). The intensity of each dot was quantified with the XDOTSREADER software (Cose) as described in (Hommais, (2001) *Mol Microbiol* **40**: 20-36). Experiments were carried out using three independent RNA preparations of TG1 planktonic flask cultures versus TG1 biofilm. For the F free TG experiment and the TG1 planktonic fermenter versus TG1 biofilm experiments, two independent RNA preparations were used. Each hybridization with each independent sample was carried out with 1 μg and 10 μg of total RNA. Comparison of the signal intensity of arrays from duplicates or from independent hybridizations showed that the results were highly reproducible (data not shown).

Statistical analysis of the Macroarray data

Genes that were statistically significantly over- and under-expressed were identified using the non-parametric Wilcoxon rank sum test. For each gene, the expression in *E. coli* TG1 flask exponential and stationary planktonic cultures (n=10 and n=12, respectively), TG flask planktonic cultures (n=4), TG1 fermenter planktonic culture (n= 4) and TG1 biofilm (n=10) or TG biofilm (n=4) were compared. Analyses were performed with one tailed tests. Genes were considered to be statistically significantly over- or under-expressed when $p<0.05$. Low (less than 0.01) or negative levels of expression were removed from the analysis.

Disruption of genes identified through macroarray analysis

fimA, *msrA*, *recA*, *cpxA* and *pspF* mutants were transferred to TG1 by P1 transduction. For the other genes, a non-polar mutation that deletes the entire target gene from the

5   initiation to the stop codon, was created by allelic exchange with the non-polar *aphA* gene cassette from Tn903. We used a 3-step PCR procedure as described in (Chaveroche, M.K., Ghigo, J.M., and d'Enfert, C. (2000) *Nucleic Acids Res* **28**: E97; Derbise, A., (2003) *FEMS Immunol Med Microbiol* **38**: 113-116) and detailed at previously.

The primers used to inactivate the 54 genes presented in this study, as well as *nlpE* and

10   *cpxR* genes, are described in Table 6.

Quantitative RT-PCR

Quantitative reverse transcription PCR (Q-RT-PCR) was used to confirm the DNA macroarray data. Total RNAs used for macroassay were used for real-time PCR and

15   RT-PCR. PCR and RT-PCR were performed using a light-cycler (Roche Diagnostics). The RNA preparation was subjected twice to DNase I (Roche Diagnostics) treatment for 30 min at room temperature to remove any contaminating genomic DNA. The enzyme was then inactivated 15 min at 65°C in the presence of 2.5 mM EDTA. Samples were checked for residual genomic DNA by real-time PCR using the cpxP-RT-5 and

20   cpxP-RT-3 primers (see Table 7). Reactions were performed in a 20 µl reaction volume using LightCycler FastStart DNA master SYBR Green I (Roche Diagnostics) according to the manufacturer's instructions. RNA samples were considered to be free of genomic DNA if no amplification was detected after at least 35 cycles of amplification. Quantitative RT-PCR reactions were performed twice with two independent RNA

25   preparations and using primers specifics for several biofilm up-regulated genes (see Table

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7) or control 16S rDNA primers (TM1, 5'-ATGACCAGCCACACTGGAAC-3' (SEQ ID NO:297) and TM2, 5'-CTTCCTCCCCGCTGAAAGTA-3' (SEQ ID NO:298)) with 50 ng of total RNA. Control 16S rDNA primers were always used to ensure the same quantity of total RNA in each reaction sample. Quantification of mRNA or 16S rRNA (as

5   control) was done using RNA master SYBR Green I (Roche Diagnostics) according to the manufacturer's instructions. Amplification of a single PCR product was confirmed by fusion curve analysis and electrophoresis on 2% agarose gels.

Construction of GFP-tagged strains

10   The strain TG1*gfp* was constructed by integration at the λ-att site of a *bla-gfp*mut3 cassette amplified from plasmid pZER1-GfpSal using a 3-step PCR procedure (Table 4S) as described in (Chaveroche, M.K., Ghigo, J.M., and d'Enfert, C. (2000) *Nucleic Acids Res* 28: E97; Derbise, A., (2003) *FEMS Immunol Med Microbiol* 38: 113-116). Plasmid pZER1-GfpSal is a gift from C.C. Guet where the *gfp*mut3 gene (Cormack *et al.*, 1996

15   *Gene* 173: 33-38) is controlled by the lambda right promoter. Strains TG1*gfpΔycfJ*, TG1*gfpΔyccA*, TG1*gfpΔcpxP* and TG1*gfpΔcpxR* were constructed by P1*vir* transduction into TG1*gfp*.

Flow chamber experiments

20   Biofilms were cultivated at 30°C in three-channel flow cells with individual channel dimensions of 1 × 4 × 40 mm. The flow system was assembled and prepared as previously described (Christensen *et al.*, 1999, *Methods Enzymol* 310: 20-42). A microscope glass cover slip (Knittel 24 × 50 mm st1; Knittel Gläser) was used as substratum for biofilm growth.

25   Inocula were prepared as follows: 16-20 h old overnight cultures in LB supplemented

with the appropriate antibiotics were harvested and resuspended in 0.9% NaCl. 250 μL of OD$_{600}$ - normalized dilutions in 0.9% NaCl (OD$_{600}$=0.05) were injected into each flow channel after medium flow was arrested. Flow was started 1 h after inoculation at a constant rate of 3 mL h^{-1} using a Watson Marlow 205S peristaltic pump.

5

Microscopy and image analysis

Biofilm development in micro-fermenters was recorded with a Nikon Coolpix 950 digital camera. Transmission and scanning laser electronic microscopy were performed on biofilm grown in micro-fermenters on thermanox slides (Nalgene) attached to the

10 internal removable glass slide and treated as described in (Prigent-Combaret *et al.*, 2000, *J Bacteriol* **181**: 5993-6002.).

For flow chamber experiments, microscopic observations and image acquisitions were performed on a Zeiss LSM510 Scanning Confocal Laser Microscope (Carl Zeiss, Jena, Germany). Images were obtained using a 40×/1.3 Plan-Neofluar oil objective.

15 Simulated three-dimensional images were generated by using the IMARIS software package (Bitplane AG, Zürich, Switzerland). Images were further processed for display using Adobe Photoshop. For COMSTAT analysis (Heydorn *et al.*, 2000, *Microbiology* **146** (**Pt 10**): 2395-2407) and quantification of the *E. coli* biofilm development with the wild type and the different mutants, each strain was grown in two separate channels, and

20 six image stacks were acquired randomly down through each channel at different time points (20 h, 45 h, 70 h and 95 h after inoculation).

25

Results

Production of mature E. coli biofilms

The capacity of different *E. coli* K12 strains to form mature biofilms was tested in M63B1-glucose minimal medium in a micro-fermenter-based continuous flow culture system (Ghigo, 2001, *Nature* 412: 442-445.). Most of the strains tested formed only thin biofilms after 2 to 5 days. However, high biomass and thick biofilm production (> 200 μM) was reproducibly achieved using *E. coli* TG1, a strain carrying the F conjugative plasmid previously shown to promote biofilm formation (Ghigo, 2001, *Nature* 412: 442-445.; Reisner *et al.*, 2003, *Mol Microbiol* 48: 933-946). To identify *E. coli* genes that are differentially expressed in mature biofilms, we compared 8 day-old TG1 biofilms to late exponential TG1 planktonic (OD= 0.6) or stationary phase cultures (OD= 3). Whereas in agitated flask and planktonic culture conditions, no surface adhesion was observed, a significant amount of contaminating biofilm formation occurred in planktonic TG1 continuous cultures grown in fermenters. This led us, in the main experiment described in this study, to compare planktonic cultures grown in agitated flasks to TG1 biofilms grown in fermenters. However, differential gene expression between planktonic and biofilm bacteria both grown in fermenters was also investigated (see discussion).

Biofilm formation has a global impact on gene expression when compared to exponential growth phase

Total RNAs were isolated from independent biofilm and exponential growth phase cultures and subjected to a stringent expression profiling procedure using *E. coli* membrane DNA macroarrays. Data were subjected to a Wilcoxon rank test. The

expression pattern and predicted function of differentially expressed genes are summarized in Fig. 1 and Fig. 7. In biofilms, 250 genes (5.8 %) were over-expressed (p< 0.05, 82% of them with p<0.005) whereas 188 genes (4.4%) were under-expressed (p<0.05, 85% of them with p<0.005). This indicates that 10.2 % of the *E. coli* genome is differentially expressed in TG1 biofilm at a statistically significant level (Fig. 1, 7 and Table 3 and 4). Among these identified genes, 1.9 % were up or down-regulated by a factor of two-fold or more.

The most significant classes of biofilm-induced genes when compared to the planktonic exponential growth phase either by level of over-expression or by number are i) genes involved in cellular processes such as envelope stress-responses (*pspABCDE, cpxP, spy, rpoE, rseA, rseB*) and stress (*recA, dinI*) as well as cell envelope biogenesis and transport (*fimA, tatE*), ii) genes involved in energy (*cyoD, sucA, sixA, nifU*) and carbohydrate metabolic functions (*rbsB, lamB*) and iii) genes of unknown function _(48 %) (Fig. 1).

The main classes of repressed genes include genes involved in amino acid, carbohydrate transport and inorganic ion transport and genes of unknown function (Fig. 7 and Table 4). In the rest of this study, we focus on genes that were found to be the most over-expressed in *E. coli* biofilms. The role and significance of the repressed functions will be reported elsewhere.

Both stationary phase and biofilm-specific genes are expressed in mature biofilms

Mature biofilms constitute heterogeneous environments where bacteria grow at different rates. This heterogeneity is proposed to be mostly dependent on nutrient availability and depth-related conditions created within the biofilm. We wished to determine to what extent the genes identified above were truly biofilm-specific or, instead,

a consequence of the stationary phase-like conditions prevailing in the mature biofilm. Total RNAs were isolated from independent stationary phase planktonic cultures, subjected to the expression profiling procedure and compared to biofilm profiling (complete comparison is published). Among the 64 genes found to be the most induced in biofilm versus exponential phase (≥two-fold ratio, see Fig. 1), 61% (39/64) of them were not induced in biofilm when compared to stationary phase (Table 1). This suggests that these 39 genes are not biofilm-specific, but may, instead, reflect the stationary phase-like growth conditions within the mature *E. coli* biofilm.

In contrast, 39% (25/64) of the remaining genes were also over-expressed in biofilm versus stationary growth phase, 24 of which with a ratio ≥2, thus defining a set of biofilm-specific genes (Table 1 and Table 5).

Validation of the macroarray data

Several approaches were used to validate the data issued from transcriptional profiling experiments. We checked the correlation between expression data and operons structure in *E. coli*. An analysis restricted to the genes with known function found to be induced by at least a two-fold factor in biofilm compared to exponentially grown cells showed that 51 % of them (21/41) were predicted to be included in 14 different operons, using the EcoCyc Database. For 10 of these 14 operons, we identified at least two members of the operon whose expression was induced in biofilms compared to exponentially grown cells. Furthermore, in order to verify the expression level changes, we then performed a Quantitative RT-PCR analysis (Q-RT-PCR) on a selection of the biofilm growth-regulated genes. Q-RT-PCR was performed for 7 of the most biofilm-induced genes compared to exponentially grown cells (*cpxP*, *ycfJ*, *ycfR*, *yebE*,

cyoD, *sucA* and *fimA*, see Fig. 1 and Table 1). Fig. 2 shows a good correlation between the data obtained by the two different techniques (r= 1.12).

These results indicate both a good internal consistency of our macroarray data as well as a good correlation between our analysis and actual mRNA level, as experimentally determined by Q-RT-PCR. To extract further functional information from our DNA-array data, we then wished to analyze the biofilm-related phenotypes of isogenic mutants of the identified biofilm-induced genes.

Functional profiling of E. coli biofilms: 20 biofilm-induced genes are involved in mature biofilm development

Among genes significantly induced in TG1 biofilms (when compared to planktonic exponential growth phase cells), 64 genes were found to be over-expressed by at least a factor of two (Table 1). To test directly the contribution of these genes to biofilm development, we deleted 23 of the 25 genes that were over-expressed in biofilms compared to both planktonic phases (biofilm-specific genes) as well as 31 of the 39 genes that were only induced in biofilms versus exponential growth phase. Mutations in *sixA*, *sucA*, *yfhN* (*nifU*), *yfhO* (*nifS*), *ybeD*, *yhhY, rpoE* and *rseA* impaired growth in M63B1 glucose minimal medium (data not shown). Mutants in these genes, along with *ftsL*, an essential cell division gene, could not be meaningfully tested for biofilm formation and were therefore excluded from further biofilm analysis. *rpoE* is an essential gene which mutations can be suppressed by extragenic mutations (De Las Penas *et al.*, 1997, *J Bacteriol* **179**: 6862-6864). Although our *rpoE* mutant did not exhibit full wild-type growth, we cannot exclude the appearance of such suppressor mutations in this mutant.

The ability to form a mature biofilm within 24 hours was assessed for each mutant and compared to TG1. Both macroscopic biofilm development in micro-fermenters and biofilm cell density after dispersion of the biofilm grown on the removable glass slide of the fermenter were examined. Twenty mutants displayed a reduced biofilm phenotype

5   (see Table 1, Fig. 3 and Fig. 8). Nine of the mutants with reduced biofilm biomass correspond to genes of known function: *fimA*, *msrA*, *rbsB*, *mdh*, *lctR*, *tatE*, *recA*, *cpxP* and *spy*.

fimA, *msrA*, *rbsB* and *mdh* are genes encoding proteins that have been already linked

10   to biofilm formation or adhesion properties (see above). As expected, adhesion appeared to be a key factor of TG1 biofilm formation. Indeed, *fimA* encodes for the major subunit of type I fimbriae, a known initial adhesion factor (Klemm and Christiansen, 1987, *Mol Gen Genet* **208**: 439-445) whose role has been previously demonstrated in biofilm formation (Austin *et al.*, 1998, *FEMS Microbiol Lett* **162**: 295-301; Cookson *et al.*, 2002,

15   *Int J Med Microbiol* **292**: 195-205; Cormio *et al.*, 1996, *Scand J Urol Nephrol* **30**: 19-24; Pratt and Kolter, 1998, *Mol Microbiol* **30**: 285-293; Watnick *et al.*, 1999, *J Bacteriol* **181**: 3606-3609). In contrast with our results, Reisner *et al.* recently showed that a *fimA* mutation had no effect on the development of biofilms formed in flow chambers by a F plasmid-bearing *E. coli* strain (Reisner *et al.*, 2003, *Mol Microbiol* **48**: 933-946).

20   Differences in strain, medium and biofilm growing system used might account for this discrepancy. *msrA* encodes a peptide methionine sulfoxide reductase (MsrA), a repair enzyme, that contributes to the maintenance of adhesins in *Streptococcus pneumoniae*, *Neisseria gonorrhoeae*, *E. coli* (Wizemann *et al.*, 1996, *Proc Natl Acad Sci U S A* **93**: 7985-7990) and in *Mycoplasma genitalium* (Dhandayuthapani *et al.*, 2001, *J Bacteriol*

183: 5645-5650), which could explain the alteration of biofilm formation in the *msrA* mutant.

The biofilm lifestyle leads to a profound modification of energy metabolism as judged by the identification of *mdh*, *rbsB* and *lctR* as biofilm-induced genes. The *rbsB* and *mdh*

5   genes have been already identified as being over-expressed in biofilms formed by pathogenic *E. coli* (Tremoulet *et al.*, 2002, *FEMS Microbiol Lett* **215**: 7-14). *rbsB* is part of the *rbsDACBK* operon that encodes high affinity transport of and chemotaxis towards D-ribose (*rbsC* and *rbsD* are also induced in biofilm, see Table 3). *mdh* encodes malate dehydrogenase, an enzyme of the TCA cycle. The *lctR* gene encodes for a regulator of

10   L-Lactate dehydrogenase. Furthermore, several sugar metabolism/transport systems are activated in biofilm (maltose transport, glycerol metabolism and uptake, galactose binding proteins, see Table 3).

Our results also suggest that mature *E. coli* biofilm formation might require Tat-dependent secretion of a specific set of proteins. Indeed, *tatE* is proposed to be

15   involved in the twin-arginine cell envelope protein transport system (Chanal *et al.*, 1998, *Mol Microbiol* **30**: 674-676). In *P. aeruginosa, tatA* and *tatB*, encoding components of this secretion system, have been shown to be induced in biofilms (Whiteley *et al.*, 2001, *Nature* **413**: 860-864), whereas *tatC* have been shown to be required for biofilm formation (Ochsner *et al.*, 2002, *Proc Natl Acad Sci U S A* **99**: 8312-8317).

20   We also observed a defect in mature biofilm formation in a *recA* mutant (Fig. 3). This underlines the importance of stress-responses in *E. coli* TG1 biofilm. Consistent with this result, several stress-response genes are over-expressed in TG1 biofilm (SOS response: *dinI, dinP, dinG, sbmC, recN, sulA*; general stress: *rpoS*; chaperones: *dnaJ* and *dnaK*; heat-shock proteins: *htpX, htpG* and *ddg*; DNA repair: *exo, xthA* and envelope stress: see

25   Table 3 and below). *cpxP* and *spy* are both linked to envelope stress response (Connolly *et*

al., 1997, *Genes Dev* **11**: 2012-2021; Danese and Silhavy, 1997, *Genes Dev* **11**: 1183-1193; Raivio and Silhavy, 2001, *Annu Rev Microbiol* **55**: 591-624) and will be investigated below.

We could also assign a biofilm-related function to 11 genes of previously unknown

5   function (*ycfJ, ycfR, yoaB, yqcC, yggN, yneA, yccA, yfcX, yghO, yceP* and *ygiB*). YfcX may be required for fatty acid utilization as a carbon source in anaerobic conditions (Campbell *et al.,* 2003, *Mol Microbiol* **47**: 793-805). Among these 11 genes, 5 encode putative extra-cytoplasmic proteins (*ycfJ, ycfR, yqcC, yneA, yccA*). YcfJ is homologous to UmoD of *P. mirabilis,* a protein that negatively regulates the *flhDC* flagellar and

10   swarming master operon (Dufour *et al.,* 1998, *Mol Microbiol* **29**: 741-751). *yccA* is a putative *cpx*-regulon member (De Wulf *et al.,* 2002, *J Biol Chem* **277**: 26652-26661) encoding a protein of unknown function but it has been shown to be a substrate for the membrane protease FtsH (Kihara *et al.,* 1998, *J Mol Biol* **279**: 175-188). Among the mutants lacking any one of these five putative extra-cytoplasmic proteins, Δ*ycfJ* and

15   Δ*yccA* were the most affected for mature biofilm formation, with a reduction of about 50% compared to wild type strain TG1 (Fig. 8).

To investigate the biofilm-related role of these two putative membrane proteins further and to confirm their importance in mature biofilm formation, we genetically introduced the Green Fluorescent Protein (GFP) gene into the wild type strain TG1, and

20   in the mutant strains TG1Δ*ycfJ* and TG1Δ*yccA*. This allowed us to compare biofilm formation between TG1*gfp* and TG1*gfp*Δ*ycfJ* and TG1*gfp*Δ*yccA* in continuous flow chamber cultures, another well established experimental model that is a non-invasive means of observing where the spatial arrangement of the cells is preserved. This experimental system allows the quantitative, real-time monitoring of biofilm architecture

25   development using Confocal Laser Scanning Microscopy and COMSTAT analysis

(Heydorn *et al.*, 2000, *Microbiology* **146** (**Pt 10**): 2395-2407) (Fig. 4). Initial adhesion of

the two *ycfJ* and *yccA* mutants was not affected, as measured by substrate coverage and

biomass analysis. However, the maturation of the biofilm formed by these two mutants

was strongly delayed, especially for the *yccA* mutant. Indeed, in the *yccA* mutant, the

5   accumulated biomass remained very low over time and typical biofilm mushroom

structures appeared only sporadically and much later compared to wild type strain TG1

(see Fig. 4). This suggests a role of YcfJ and YccA proteins in biofilm maturation.

These results demonstrate the involvement in mature biofilm formation of 30% of

the most highly expressed genes identified in our study. 50 % of these genes (10/20) were

10   induced in biofilm versus both exponential and stationary growth phase (*cpxP*, *spy*, *tatE*,

lctR, *mdh*, *rbsB*, *ygiB*, *yqcC*, *yceP* and *yfcX*) whereas the other 50 % (10/20) were only

induced in biofilm versus exponential growth phase (*fimA*, *msrA*, *recA*, *yoaB*, *ycfJ*, *ycfR*,

yneA, *yccA*, *yggN* and *yghO*) (see Table 1 and Table 5).

15   *Biofilm-induced genes are not involved in the early stage of biofilm formation*

A failure to form a wild type mature biofilm could result from an initial adhesion

defect. Therefore, we investigated whether the genes identified as over-expressed in

mature TG1 biofilms and that impaired mature biofilm formation when mutated were also

involved in the early adhesion steps. For this, we tested this mutants in a static microtiter

20   plate-based assay that has been widely used to study the first steps of biofilm formation

(Genevaux *et al.*, 1996, *FEMS Microbiol Lett* **142**: 27-30; O'Toole *et al.*, 1999, *Methods*

Enzymol **310**: 91-109). With the exception of *fimA*, the early adhesion capacity of the

mutants could not be distinguished from the parental strain (Fig. 9). This result indicates

that most genes over-expressed in mature biofilms are not involved in the early steps of

25   this process and confirms that they participate in mature biofilm functions.

Comparison of E. coli *F⁺/F⁻ biofilm global response: general relevance to* E. coli

biofilm

In this study, we used an *E. coli* strain carrying a conjugative plasmid, a widespread

5   situation which promotes biofilm formation (Ghigo, 2001, *Nature* **412**: 442-445; Reisner

et al., 2003., *Mol Microbiol* **48**: 933-946). To distinguish general features of *E. coli*

biofilms from those specific to our model, we analyzed the transcription profile of the *E.*

coli strain TG, an F-free isogenic derivative of TG1. This control is of particular

relevance because some of the genes found to be the most over-expressed (*pspA, cpxP*)

10   have either been shown to be related to the conjugation process (*cpx* stands for

conjugation plasmid expression (McEwen and Silverman, 1980, *Proc Natl Acad Sci U S*

A **77**: 513-517) or to stress-responses that could correlate with the expression of

membrane appendages such as conjugative pili. TG forms a thin and fragile biofilm after

5 days of culture in micro-fermenters (data not shown). Total RNA was isolated from *E.*

15   *coli* TG biofilm and flask planktonic exponential cultures, and was subjected to the same

macroarray analysis as described for TG1. TG1 and TG biofilms were not strictly

comparable in terms of depth and structure (and therefore, possibly, for biofilm-induced

responses). As expected, some functions induced in TG1, for instance RecA and part of

the SOS stress pathway, were not induced in TG (Table 1), suggesting that F-specific,

20   possibly transfer-related, responses are induced in TG1 biofilm. Despite this fact, 33% of

the genes induced in TG1 biofilm by an over two-fold factor were also found to be

statistically significantly over-expressed in TG biofilm (including *cpxP, rseA, rseB, spy,*

psp operon members, *tatE,* and *fimA,* see Table 1). This demonstrates that many of the

biofilm-induced genes identified in this study are F-independent and part of a general *E.*

25   *coli* K12 biofilm response.

28

Envelope stress pathways in *E. coli* mature biofilm

cpxP is one the most over-expressed genes in *E. coli* TG1 biofilms versus planktonic

growth phase (Fig. 1, Table 1 and Table 5). *cpxP* is a target of the *cpx* two-component

system, which is known to respond to a variety of extra-cytoplasmic stress (envelope

stress) (Raivio and Silhavy, 2001, *Annu Rev Microbiol* 55: 591-624).

We therefore investigated the effect of deletion mutations in key components of the

cpx pathway on biofilm formation. As shown in Figure 5, inactivation of the

sensor-regulator components of the *cpx* system (*cpxA*, *cpxR*), but also of *cpxP* and of *nlpE*

affected biofilm formation in micro-fermenters. A mutation in *spy* (a biofilm-induced

cpxP homolog) has no effect on biofilm biomass. *rpoE* and *rseA* mutants displayed a

growth rate defect and consequently could not be studied in micro-fermenters. A

mutation in *rseB*, the second anti-sigma E factor of the RpoE envelope stress pathway,

did not affect growth and a *rseB* mutant formed a wild type biofilm. Whereas it is difficult

to conclude that the *rpoE* pathway has a role in biofilm formation, the *cpx* pathway

appears to contribute to biofilm development, based on the morphological effects caused

by mutations in several of its key components. Indeed, the biofilms produced by both

TG1Δ*cpxR* and TG1Δ*cpxP* in micro-fermenters were very fragile compared to wild type

TG1 biofilms. TG1Δ*cpxP* biofilm was made of large plaques, in strong contrast to the

homogeneous TG1 biofilm (Fig. 6ABC). Consistent with this observation, a detailed

electron microscopy analysis revealed that a *cpxP* mutation strongly altered biofilm

macromorphology (Fig. 6DE). Despite its fragility, no clear structural defect could be

detected in the TG1Δ*cpxR* biofilm (data not shown). Even though slight structural

differences could also be seen in the TG1Δ*spy* mutant biofilms, structural alterations were

not found in *nlpE*, *cpxA* nor *rseB* mutant biofilms grown in micro-fermenters (data not shown).

To further investigate the role of *cpxP* and *cpxR*, we introduced a *gfp* allele into TG1Δ*cpxP* and TG1Δ*cpxR* and we compared their biofilm formation to the parental TG1*gfp* strain in continuous flow chamber cultures. Single cells and very small colonies were observed on the surface for these two mutants during the initial steps of biofilm development in contrast to the wild-type that forms normal three-dimensional colonies (Fig. 4, 20 and 45 h). Furthermore, both *cpxP* and *cpxR* mutants were also strongly affected for maturation of the biofilm (Fig. 4). These experiments suggest that stress envelope pathways are involved in the establishment of a structured mature biofilm in *E. coli*.

Phage-shock protein operon (*psp*) is expressed in response to a variety of environmental and intracellular stresses including processes related to protein insertion in the outer membrane (Weiner and Model, 1994, *Proc Natl Acad Sci U S A* **91**: 2191-2195). While the precise functions of the *psp* genes are not understood, they help to ensure survival of *E. coli* in adverse conditions, suggesting that *psp* genes are part of a stress-response operon (Model *et al.*, 1997, *Mol Microbiol* **24**: 255-261). In our analysis, *pspA* and other members of the operon (*pspBCDE*) were consistently over-expressed in biofilm (Fig. 1, Table 3 and Table 5). Nevertheless, the disruption of the *pspABCDE* operon did not have a major impact on early (Fig. 9) or late biofilm formation nor on biofilm structure (data not shown).

Discussion

In this study we investigated the differences in gene expression between *E. coli* K12 mature biofilm and planktonic laboratory cultures. Using DNA macroarrays we showed that the biofilm lifestyle, while sharing similarities with the stationary growth phase, triggers the expression of specific sets of genes.

Modifications of E. coli *K12 gene expression induce by the biofilm lifestyle*

The use of large scale fusion technology had already suggested that a significant fraction of the bacterial genome could be involved in biofilm physiology (Prigent-Combaret *et al.*, 1999, *J Bacteriol* **181**: 5993-6002). Accordingly, *P. putida* and *P. aeruginosa* biofilm proteome analyses showed that a large number of genes are differentially regulated during biofilm development (Sauer and Camper, 2001, *J Bacteriol* **183**: 6579-6589; Sauer *et al.*, 2002, *J Bacteriol* **184**: 1140-1154). In contrast, a transciption profiling of the *P. aeruginosa* planktonic and biofilm phases led to the conclusion that only 1% of *P. aeruginosa* genes display over a two-fold difference in gene expression (Whiteley *et al.*, 2001, *Nature* **413**: 860-864).

In *E. coli*, Schembri *et al.* recently showed that approximately 5 to 10 % of the *E. coli* genes exhibited altered microarray expression profiles when compared planktonic growth phases and young biofilm cultures. They hypothesized that this could be due to the rather early stages of biofilm development analyzed in their study, where the still ongoing switch from planktonic to sessile growth could result in a high level of transient gene expression (Schembri *et al.*, 2003, *Mol Microbiol* **48**: 253-267).

Here, we compared mature biofilms to the planktonic exponential growth phase and showed that, as in the case of mature *P. aeruginosa* biofilms, only a small fraction (1.9 %)

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of the *E. coli* genes are differentially expressed by more than a factor of two. However,
below that threshold, biofilm formation still leads to the statistically significant
differential expression of more than 10% of the *E. coli* genome. These results therefore
support the proposal that biofilm formation results in and from significant differences in

5   the overall make-up of bacterial cells (Sauer, 2003, **Genome Biol** 4: **219**; Stoodley *et al.*,
2002, **Annu Rev Microbiol** 56: **187-209**).

Mature biofilm cells have been proposed to have stationary growth phase traits such
as reduced growth and metabolic activity. To investigate the stationary phase character of
bacterial life within biofilm, we also compared the expression pattern of stationary phase

10   cultures with those determined for the exponential growth phase and the mature biofilm.
Biofilm-specific genes, *i.e.* genes differentially regulated in biofilm versus both forms of
planktonic phases, correspond to 4% of the genome (118 over- and 53 under-expressed/
4290) and this proportion decreases to less than 1% (0.67%, 23 over and 6 under/4290)
for genes varying by a factor of more than two. When one only considers the genes

15   induced in response to the stationary growth phase character of the biofilm lifestyle, these
genes represent 3% of the genome. The biofilm lifestyle, while sharing similarities with
the stationary growth phase, thus triggers the expression of specific sets of genes.

Functional profiling of the biofilm-induced genes

20   The biological importance of the differential gene expression exhibited upon biofilm
versus planktonic growth was tested by the disruption of the majority of the
highly-induced genes in biofilms, including all biofilm-specific induced genes. We show
that, while the mutants were not impaired in initial steps of adhesion to surfaces (with the
exception of *fimA*), a third of them (20 genes) were affected in the biofilm maturation

25   (Table 1, Fig. 3 and Fig. 8). This high proportion of genes involved in the biofilm

maturation strongly supports the pertinence of our analysis. Among these 20 genes, half correspond to biofilm-specific genes whereas the other half was only induced in biofilms versus exponential growth phase (see Table 1 and Table 5). This indicates that the development of a full mature biofilm requires not only biofilm-specific genes but also

5   genes related to the stationary phase character of the biofilm. The individual role of some of these newly identified genes is currently being investigated.

Biofilm-related physiological functions

We show that genes found to be the most over-expressed in TG1 biofilm versus

10   exponential growth phase were also part of the *E. coli* F-free biofilm response, therefore indicating that genes identified in this study are involved in the general response developed in mature *E. coli* K12 biofilms. Those genes are not distributed randomly into all potential functional classes. Instead they display a strong bias toward specific functional-categories and we propose that they are part of the biofilm genetic signature.

15   Genes whose expression is required for full maturation of TG1 biofilm belong to functions linked to adhesion (*fimA*, *msrA*), energy metabolism (*rbsB*, *mdh*, *lctR*), transport (*tatE*), general stress (*recA*), and envelope stress response (*cpxP* and *spy*). However, it is likely that many genes identified in our study are not specifically involved in biofilm-specific functions but rather correspond to adaptive responses to the biofilm

20   environment. Mutations in many biofilm-induced genes that also correspond to information storage and processing, metabolism, cellular processes and unknown functions have indeed no effect on TG1 biofilm formation (Table 1).

Moreover, 48% of the genes significantly over-expressed in biofilms versus exponential growth phase were of uncharacterized function. Compared to 19.6% of such

25   genes found in the *E. coli* genome (Serres *et al.*, 2001, *Genome Biol* **2**: RESEARCH0035),

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this high proportion of genes of unknown functions expressed in mature biofilm suggests that new aspects of *E. coli* biology are adopted during biofilm formation. We show that 11 of these uncharacterized genes are necessary for full mature biofilm formation, thus experimentally assigning them a biofilm-related function (Table 1, Fig. 3 and Fig. 8).

5   Among them 5 encode putative membrane proteins that could be of particular relevance when considering the importance of envelope-related physiology within a biofilm.

Consistent with the drastic phenotypic changes occurring inside biofilms, we found that 15 % of the genes identified as over- or under-expressed in biofilms versus exponential growth phase are involved in either energy processes or carbohydrate

10   metabolism (Fig. 1, Table 1 and 3). Despite the presence of polysaccharides in the TG1 biofilm (data not shown), we could not clearly associate the expression of any of those genes with the production of the biofilm matrix (*i.e.*, cellulose, colanic acid). This could reflect, among other explanations, a lack of sensitivity of our approach due to the averaging occurring while extracting transcription information from the heterogeneous

15   bacterial biofilm population.

A partial comparison of the most over-expressed genes in our analysis (>2 fold factor) and in the study by Schembri *et al.* (> 8 fold factor) only revealed a few genes identified as over-expressed in *E. coli* biofilm in both studies (*rbsB, b0836, yfjO, yceP, glgS, ydeW, yneA, yqeC, ylcC, rplV, rplD, rpsS, b1550, rplP, rpsR, flu, rplM, ppc, oppA,*

20   *gatD, cydA, atpB, rpsN, malK, atpG*) (Schembri *et al.*, 2003, *Mol Microbiol* **48**: 253-267). Three of these genes (*rbsB, yceP, yneA*) were nevertheless also found here to be required for mature biofilm formation. This relatively low overlap between the two studies may be due to technical differences. Different scenarios were used in terms of strain background, media and experimental set-up. This could also reflect the difference in the gene

25   expression pattern between two biofilms at very different stages of maturation (*i.e.* young

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and thin biofilms in Schembri *et al.* versus mature and thick biofilms in our study). Further studies comparing the expression profile of *E. coli* biofilms at different maturation stages within the same experimental set-up will provide a more dynamic view of biofilm gene expression.

5 *Heterogeneity of oxygen conditions in* E. coli *K12 biofilms*

Biofilms are heterogeneous environments and, with respect to aerobiosis, our analysis supports these results. In the main experiment described in this study, we compared exponentially grown agitated flask cultures to TG1 biofilm in aerated conditions. Under

10 these conditions, numerous genes known to be induced by aerobiosis were also induced in biofilms, including some genes for TCA cycle enzymes (e.g. *aceB*, *cyo* operon *members, fadB, mdh, glpD, sucAB*). In addition, some genes known to be repressed by aerobiosis were repressed in biofilms (eg. *adhE, cydAB, dcuC, focA, fumB*). This tends to indicate that our biofilms were mainly grown under aerobic conditions. Consequently, we also

15 compared differential gene expression between TG1 biofilms and TG1 planktonic cultures, both grown in aerated fermenters (data not shown). In this configuration, we clearly observed that some typical aerobic genes were induced in biofilms whereas others were repressed. This was also the case for typical anaerobic genes. This could reflect the heterogeneity of the aerobic conditions in biofilms, in which external bacteria are in

20 contact with oxygen while internal bacteria are in conditions close to anaerobiosis.

Stress-responses in biofilms

Our study revealed that a major physiological response to biofilm formation is the induction of stress-responses. Interestingly, such a stress-response induction may also

25 take place in *P. aeruginosa* biofilms. Indeed, the most highly activated genes identified in a *P. aeruginosa* biofilm transcriptome analysis were those of temperate bacteriophages

(Whiteley et al., 2001, Nature **413**: 860-864). As stresses are known to induce prophages and other mobile genetic elements, our results suggest that *Pseudomonas* prophage induction may be a consequence of stresses created by the drastic conditions that prevail inside the biofilm. As such, stress may well be a key factor in the mechanisms that lead to

5   the observed antibiotic resistance inside biofilm communities.

Owing to the possible role of cell-cell and cell-surface interactions in biofilm, it may be of significance that envelope stress genes such as *cpxP*, *spy* and the *psp* genes are consistently induced in this environment. CpxP may inhibit the *cpx*-mediated induction through a direct interaction with the two-component system sensor CpxA, while Spy may

10   play a similar role on the *rpoE* pathway (Raivio et al., 2000, *Mol Microbiol* **37**: 1186-1197). The *cpx* system is known to respond to envelope stresses such as over-production and misfolding of membrane proteins or elevated pH (Raivio and Silhavy, 2001, *Annu Rev Microbiol* **55**: 591-624). However, relatively little is known about the physiological role of envelope stress-responses. Recently, adhesion of *E. coli*

15   cells to hydrophobic but not hydrophilic surfaces was shown to activate the *cpx* system, including *cpxP*, through a process called surface sensing which requires both *cpxR* and *nlpE* (Otto and Silhavy, 2002, *Proc Natl Acad Sci U S A* **99**: 2287-2292). Consistently, we find that *cpxP* and *spy* are highly induced in mature biofilms where bacteria are *de facto* in contact with the hydrophobic surfaces of other cells.

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Our results thus provide additional experimental evidence that stress response pathways are key factors in biofilm formation. The structure of biofilms grown in micro-fermenters is altered in a *cpxP* mutant (Fig. 6) and to a lesser extent in a *spy* mutant. Observation of *spy* mutant biofilms by transmission electron microscopy also revealed a

25   high proportion of spheroblasts as compared to wt TG1 (data not shown), suggesting a

possible cause for the affected structure of the biofilm in this mutant. In addition, a *cpxP* and a *cpxR* mutant are both impaired in forming wild type micro-colonies (Fig. 4). This strongly corroborates the idea that *cpxP* and *cpxR* mutants have reduced cell-to-cell adherence, since any growth up in the water column will be counteracted by the shearing forces of the flow. It appears, then, that the inappropriate expression of the *cpx* regulated genes in biofilm, *i.e.* a derepression of the *cpx* regulon in the *cpxP* mutant or an absence of induction of the *cpx* regulon in the *cpxR* mutant, leads to an alteration of the process of biofilm formation. Considering the importance of environmental conditions in biofilm formation, two component systems, which sense perturbations or changes in the bacterial environment, might play a regulatory role in bacterial biofilm formation, a proposal that requires further investigation.

Our analysis identified the biofilm mode of growth as an environment that induces the expression of the *pspABCDE* stress operon. However no biofilm-related phenotype could be observed in a strain deleted for the *pspABCDE* operon. Nevertheless, the deletion of *pspF*, a constitutively expressed positive regulator of the *pspABCDE* operon, affects biofilm formation (Fig. 8). Since *pspABCDE* is not required for biofilm formation, *pspF* might also regulate a biofilm-related locus that is not part of *pspABCDE* operon. Evidence for such an additional PspF regulated target has been provided in the case of *Yersinia enterolitica psp* regulon (Darwin and Miller, 2001, *Mol Microbiol* **39**: 429-444).

Changes in gene expression and biofilm development

The changes in gene expression demonstrated here and in other studies could be considered either as part of the *E. coli* biofilm development (needed for maturation) or as caused by the conditions progressively created within the biofilm during its maturation (consequence of the maturation). The first hypothesis implies that the biofilm formation

is a developmental process in which genetic checkpoints could control the maturation of the biofilm by inducing a succession of biofilm-specific genes. Whereas 8 mutations out of 54 mutants created in this study display a 50% decrease in biofilm biomass and maturation, none of them lead to a total loss of biofilm formation. Considering the

5   existence of multiple and partially overlapping or complementing pathways that can lead to biofilm formation, this result, without formally excluding the existence of a biofilm developmental program, rather speaks in favor of the second working hypothesis. In this case, most changes observed in biofilm gene induction could be a consequence of, rather than a prerequisite for the biofilm maturation.

10

The results presented here provide new insights into the global effect triggered by biofilm formation in *E. coli*. By monitoring the changes in gene expression occurring in mature biofilms, we have identified biofilm-related physiological pathways and previously-uncharacterized biofilm-induced genes. This may lead to new biofilm control

15   strategies that will likely hinge upon a better understanding of biofilm-induced physiological responses.

Discussion

20   In this study we investigated the differences in gene expression between *E. coli* K12 mature biofilm and planktonic laboratory cultures. Using DNA macroarrays we showed that the biofilm lifestyle, while sharing similarities with the stationary growth phase, triggers the expression of specific sets of genes.

25   *Modifications of* E. coli *K12 gene expression induce by the biofilm lifestyle*

38

The use of large scale fusion technology had already suggested that a significant fraction of the bacterial genome could be involved in biofilm physiology (Prigent-Combaret et al., 1999, J Bacteriol 181: 5993-6002). Accordingly, P. putida and P. aeruginosa biofilm proteome analyses showed that a large number of genes are differentially regulated during biofilm development (Sauer and Camper, 2001, J Bacteriol 183: 6579-6589; Sauer et al., 2002, J Bacteriol 184: 1140-1154). In contrast, a transcription profiling of the P. aeruginosa planktonic and biofilm phases led to the conclusion that only 1% of P. aeruginosa genes display over a two-fold difference in gene expression (Whiteley et al., 2001, Nature 413: 860-864).

In E. coli, Shembri et al. recently showed that approximately 5 to 10 % of the E. coli genes exhibited altered microarray expression profiles when compared planktonic growth phases and young biofilm cultures. They hypothesized that this could be due to the rather early stages of biofilm development analyzed in their study, where the still ongoing switch from planktonic to sessile growth could result in a high level of transient gene expression (Schembri et al., 2003, Mol Microbiol 48: 253-267).

Here, we compared mature biofilms to the planktonic exponential growth phase and showed that, as in the case of mature P. aeruginosa biofilms, only a small fraction (1.9 %) of the E. coli genes are differentially expressed by more than a factor of two. However, below that threshold, biofilm formation still leads to the statistically significant differential expression of more than 10% of the E. coli genome. These results therefore support the proposal that biofilm formation results in and from significant differences in the overall make-up of bacterial cells (Sauer, 2003, Genome Biol 4: 219; Stoodley et al., 2002, Annu Rev Microbiol 56: 187-209).

Mature biofilm cells have been proposed to have stationary growth phase traits such as reduced growth and metabolic activity. To investigate the stationary phase character of bacterial life within biofilm, we also compared the expression pattern of stationary phase cultures with those determined for the exponential growth phase and the mature biofilm.

5 Biofilm-specific genes, *i.e.* genes differentially regulated in biofilm versus both forms of planktonic phases, correspond to 4% of the genome (118 over- and 53 under-expressed/ 4290) and this proportion decreases to less than 1% (0.67%, 23 over and 6 under/4290) for genes varying by a factor of more than two. When one only considers the genes induced in response to the stationary growth phase character of the biofilm lifestyle, these

10 genes represent 3% of the genome. The biofilm lifestyle, while sharing similarities with the stationary growth phase, thus triggers the expression of specific sets of genes.

Functional profiling of the biofilm-induced genes

The biological importance of the differential gene expression exhibited upon biofilm

15 versus planktonic growth was tested by the disruption of the majority of the highly-induced genes in biofilms, including all biofilm-specific induced genes. We show that, while the mutants were not impaired in initial steps of adhesion to surfaces (with the exception of *fimA*), a third of them (20 genes) were affected in the biofilm maturation (Table 1, Fig. 3 and Fig. 8). This high proportion of genes involved in the biofilm

20 maturation strongly supports the pertinence of our analysis. Among these 20 genes, half correspond to biofilm-specific genes whereas the other half was only induced in biofilms versus exponential growth phase (see Table 1 and Table 5). This indicates that the development of a full mature biofilm requires not only biofilm-specific genes but also genes related to the stationary phase character of the biofilm. The individual role of some

25 of these newly identified genes is currently being investigated.

40

Biofilm-related physiological functions

We show that genes found to be the most over-expressed in TG1 biofilm versus exponential growth phase were also part of the *E. coli* F-free biofilm response, therefore indicating that genes identified in this study are involved in the general response developed in mature *E. coli* K12 biofilms. Those genes are not distributed randomly into all potential functional classes. Instead they display a strong bias toward specific functional categories and we propose that they are part of the biofilm genetic signature. Genes whose expression is required for full maturation of TG1 biofilm belong to functions linked to adhesion (*fimA, msrA*), energy metabolism (*rbsB, mdh, lctR*), transport (*tatE*), general stress (*recA*), and envelope stress response (*cpxP* and *spy*). However, it is likely that many genes identified in our study are not specifically involved in biofilm-specific functions but rather correspond to adaptive responses to the biofilm environment. Mutations in many biofilm-induced genes that also correspond to information storage and processing, metabolism, cellular processes and unknown functions have indeed no effect on TG1 biofilm formation (Table 1).

Moreover, 48% of the genes significantly over-expressed in biofilms versus exponential growth phase were of uncharacterized function. Compared to 19.6% of such genes found in the *E. coli* genome (Serres *et al.*, 2001, *Genome Biol* 2: RESEARCH0035), this high proportion of genes of unknown functions expressed in mature biofilm suggests that new aspects of *E. coli* biology are adopted during biofilm formation. We show that 11 of these uncharacterized genes are necessary for full mature biofilm formation, thus experimentally assigning them a biofilm-related function (Table 1, Fig. 3 and Fig. 8). Among them 5 encode putative membrane proteins that could be of particular relevance when considering the importance of envelope-related physiology within a biofilm.

Consistent with the drastic phenotypic changes occurring inside biofilms, we found that 15 % of the genes identified as over- or under-expressed in biofilms versus exponential growth phase are involved in either energy processes or carbohydrate metabolism (Fig. 1, Table 1 and 3). Despite the presence of polysaccharides in the TG1

5   biofilm (data not shown), we could not clearly associate the expression of any of those genes with the production of the biofilm matrix (*i.e.*, cellulose, colanic acid). This could reflect, among other explanations, a lack of sensitivity of our approach due to the averaging occurring while extracting transcription information from the heterogeneous bacterial biofilm population.

10   A partial comparison of the most over-expressed genes in our analysis (>2 fold factor) and in the study by Schembri *et al.* (> 8 fold factor) only revealed a few genes identified as over-expressed in *E. coli* biofilm in both studies (*rbsB, b0836, yfjO, yceP, glgS, ydeW, yneA, yqeC, ylcC, rplV, rplD, rpsS, b1550, rplP, rpsR, flu, rplM, ppc, oppA, gatD, cydA, atpB, rpsN, malK, atpG*) (Schembri *et al.*, 2003, *Mol Microbiol* **48**: 253-267).

15   Three of these genes (*rbsB, yceP, yneA*) were nevertheless also found here to be required for mature biofilm formation. This relatively low overlap between the two studies may be due to technical differences. Different scenarios were used in terms of strain background, media and experimental set-up. This could also reflect the difference in the gene expression pattern between two biofilms at very different stages of maturation (*i.e.* young

20   and thin biofilms in Schembri *et al.* versus mature and thick biofilms in our study). Further studies comparing the expression profile of *E. coli* biofilms at different maturation stages within the same experimental set-up will provide a more dynamic view of biofilm gene expression.

25   *Heterogeneity of oxygen conditions in* E. coli *K12 biofilms*

Biofilms are heterogeneous environments and, with respect to aerobiosis, our analysis supports these results. In the main experiment described in this study, we compared exponentially grown agitated flask cultures to TG1 biofilm in aerated conditions. Under these conditions, numerous genes known to be induced by aerobiosis were also induced in

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10 cultures, both grown in aerated fermenters (data not shown). In this configuration, we clearly observed that some typical aerobic genes were induced in biofilms whereas others were repressed. This was also the case for typical anaerobic genes. This could reflect the heterogeneity of the aerobic conditions in biofilms, in which external bacteria are in contact-with oxygen while internal bacteria are in conditions close to anaerobiosis.

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20 a *P. aeruginosa* biofilm transcriptome analysis were those of temperate bacteriophages (Whiteley *et al.*, 2001, *Nature* 413: 860-864). As stresses are known to induce prophages and other mobile genetic elements, our results suggest that *Pseudomonas* prophage induction may be a consequence of stresses created by the drastic conditions that prevail inside the biofilm. As such, stress may well be a key factor in the mechanisms that lead to

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10   strategies that will likely hinge upon a better understanding of biofilm-induced physiological responses.

Genes a	b	Ratio c	Rank d	Phenotype e	TG f	Function - description g
INFORMATION STORAGE AND PROCESSING						
J : Translation, ribosomal structure and metabolism						
rplY*	b2185	2.23	52	ND		50S ribosomal subunit protein L25
rne#	b1084	2.06	58	ND		RNase E, mRNA turnover, maturation 5S RNA
K : Transcription						
È lctR#	b3604	4.76	14	-		Regulator of L-Lactate dehydrogenase genes
L : DNA replication, recombination and repair						
È recA*	b2699	2.30	51	-		DNA strand exchange and renaturation. SOS
dinI#	b1061	2.02	61	wt		Inhibits RecA-mediated self-cleavage. SOS
METABOLISM						
C : Energy production and conversion						
cyoD*	b0429	7.41	3	wt		Cytochrome o oxidase subunit IV
sucA*	b0726	6.54	7	NA		2-oxoglutarate dehydrogenase
fdhF*	b4079	3.85	23	wt		S subunit of formate dehydrogenase H
cyoC*	b0430	3.59	26	wt		Cytochrome o oxidase subunit III
nifU*	b2529	3.41	27	NA		Formation of [fe-s] clusters in iron-sulfur proteins
sixA#	b2340	2.74	41	wt	√	Phosphatase affecting ArcB phosphorelay
sucD*	b0729	2.69	43	NA		Succinyl-CoA synthetase, alpha subunit
glpQ#	b2239	2.50	46	ND		Glycerol-3-phosphate diesterase, periplasmic
È mdh#	b3236	2.19	53	-		Malate dehydrogenase
G : Carbohydrate transport and metabolism						
lamB#	b4036	2.94	36	wt		Phage lambda receptor, maltose receptor
È rbsB#	b3751	2.41	48	-		D-ribose periplasmic binding protein, chemotaxis
sfsA*	b0146	1.97	64	ND		Regulatory protein for maltose metabolism
E : Amino acid transport and metabolism						
gadA#	b3517	3.15	30	wt	√	Glutamate decarboxylase isozyme
nifS*	b2530	1.98	62	NA		Cysteine desulfurase
I : Lipid metabolism						
fadB*	b3846	4.18	20	wt		Fatty acid oxidation complex; 4-enzyme protein
Q : Secondary metabolites biosynthesis, transport and metabolism						
ucpA*	b2426	2.32	50	ND		Oxido reductase, dehydrogenase/reductase family
CELLULAR PROCESSES						
D : Cell division and chromosomal partitionning						
ftsL*	b0083	4.34	17	NA		Cell division and growth, septum localization
sulA*	b0958	3.07	33	wt	47	Inhibits cell division. SOS
O : Post-translational modification, protein turnover, chaperones						
eco*	b2209	6.17	9	wt		Ecotin, a periplasmic serine protease inhibitor

Table 1. Over-expressed genes (≥2) in *E. coli* TG1 and TG biofilms versus exponential growth phase

a : Gene names according to *E. coli* Colibri database.

5   b : Gene names according to Blattner nomenclature.

c : Ratio of gene expression in *E. coli* biofilm versus gene expression in planktonic cultures.

d : Rank position ;1= the most over-expressed gene in *E. coli* biofilm.

e : Biofilm phenotype of the mutants : ND : not determined ; NA : not applicable due to

10   growth defect in M63B1 glucose medium ; wt : similar to wild type ; - : biofilm reduced compared to wt ; Struct : biofilm structure impaired compared to wt.

f : √ genes also found to be significantly over-expressed in F minus *E. coli* strain TG.

g : Function description according to *E. coli* Colibri database.

h : *pspF* was expressed by only a 1.22 factor in TG1 biofilm but has been included for

15   comparison with other members of the *psp* operon.

Arrow: mutants affected for biofilm formation.

*: genes that were not induced in TG1 biofilm versus stationary phase.

#: genes that were also induced in TG1 biofilm versus stationary phase by at least a factor of two. These genes are also summarized in Table 3S.

20   The genes have been classified according to the COGs functional categories annotation system used by the NCBI.

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Table 2. Strains and plasmids used in this study

* Additional individual mutants in the following genes: *cutC, cyoC, dinI, eco, fadB, fdhF, gadA, lctR, malM-G, mdh, nifS, nifU, nlpE, pspA-E, rbsB, rpoE, rseB, sixA, sodC, spy, sucA, sulA, tatE, ybeD, ybjF, yccA, yceP, ycfJ, ycfL, ycfR, ydcI, yebE, yfcX, yggN, yghO, ygiB, yhhY, yiaH, yjbO, yneA, yoaB, yqcC, yqeC*, were named TG1Δ☐*gene.name*]::*aphA*, (Km^R).

Strain / plasmid	Relevant characteristics	Reference / source
***E. coli* strains**		
PAP6181	K1519*pspF::miniTn10* (Tet^R)	(Jovanovic *et al.*, 1996)
PHL904	*cpxA*:: Ω*cat* (Cm^R)	(Dorel *et al.*, 1999)
RG075	MG1655Δ*msrA*::Ω*Spec* (Spec^R)	A gift of F. Barras
STC27	*fimA1::cat* (Cm^R)	(Pratt and Kolter, 1998)
TG1	F'[*traD36 proAB+ lacI^q lacZ*ΔM15] *supE hsd*Δ5 *thi* Δ(*lac-proAB*)	Laboratory collection
TG	A F minus derivative of TG1	Laboratory collection
TG1Δ*cpxA*	TG1*cpxA*:: Ω*cat* (Cm^R)	This work
TG1Δ*cpxP*	TG1Δ*cpxP*::Δfrt	This work
TG1Δ*cpxR*	TG1Δ*cpxR*::Δfrt	This work
TG1Δ*fimA*	TG1Δ*fimA*::*cat* (Cm^R)	This work
TG1Δ*msrA*	TG1Δ*msrA*::Ω*Spec*, (Spec^R)	This work
TG1Δ*pspF*	TG1*pspF::miniTn10* (Tet^R)	This work
TG1*recA*	TG1*recA56 SrlC300::Tn10* (Tet^R)	Laboratory collection
TG1Δ*rseA*	TG1Δ*rseA*::Δfrt	This work
TG1*gfp*	TG1λatt::*gfp-bla*, (Amp^R)	A gift of A. Roux
TG1*gfp*Δ*cpxP*	TG1Δ*cpxP* λatt::*gfp-bla* (Amp^R)	This work
TG1*gfp*Δ*cpxR*	TG1Δ*cpxR* λatt::*gfp-bla* (Amp^R)	This work
TG1*gfp*Δ*yccA*	TG1Δ*yccA* λatt::*gfp-bla* (Km^R, Amp^R)	This work
TG1*gfp*Δ*ycfJ* *	TG1Δ*ycfJ* λatt::*gfp-bla* (Km^R, Amp^R)	This work
Plasmids		
pKOBEG	pSC101 *ts* (replicates at 30°C), *araC* arabinose-inducible λ *red*γβα operon, (Cm^R)	(Chaveroche *et al.*, 2000)
pCP20	*ts* (replicates at 30°C) plasmid bearing the *flp* recombinase gene, (Cm^R and Amp^R)	(Cherepanov and Wackernagel, 1995)

* Additional individual mutants in the following genes: *cutC, cyoC, dinI, eco, fadB, fdhF, gadA, lctR, malM-G, mdh, nifS, nifU, nlpE, pspA-E, rbsB, rpoE, rseB, sixA, sodC, spy, sucA, sulA, tatE, ybeD, ybjF, yccA, yceP, ycfJ, ycfL, ycfR, ydcI, yebE, yfcX, yggN, yghO, ygiB, yhhY, yiaH, yjbO, yneA, yoaB, yqcC, yqeC*, were named TG1Δ☐*gene.name*]::*aphA*, (Km^R).

Table 3. Genes over-expressed in *E. coli* TG1 biofilm versus exponential growth phase.

5 The genes found to be over-expressed at a significant level (P-value ≤0.05) are indicated. They have been classified according to the COGs functional categories annotation system.

a : Gene names according to *E. coli* Colibri database.

b : Gene names according to Blattner nomenclature.

10 c : Ranking position 1= the most over-expressed gene in *E. coli* biofilm.

d : Ratio of gene expression in *E. coli* biofilm versus gene expression in planktonic cultures.

e : Function description according to *E. coli* Colibri database.

Arrow: mutants affected for biofilm formation.

15

Genes		Rank	Bio/Exp	Function – description
a	b	c	d	e
INFORMATION STORAGE AND PROCESSING				
J : Translation. ribosomal structure and metabolism				
lysU	b4129	146	1.45	Lysine tRNA synthetase
miaA	b4171	79	1.84	Delta(2)-isopentenylpyrophosphate tRNA-adenosine transferase
rluC	b1086	166	1.35	Ribosomal large subunit pseudouridine synthase C
rne	b1084	58	2.06	RNase E
rplY	b2185	52	2.23	50S ribosomal subunit protein L25
K : Transcription				
crl	b0240	81	1.83	Transcriptional regulator of genes for curli
cspD	b0880	135	1.50	Cold shock protein
dniR	b0211	199	1.24	Transcriptional regulator for nitrite reductase
fruR	b0080	133	1.50	Transcriptional repressor of *fru* operon and others
idnR	b4264	227	1.18	L-idonate transcriptional regulator
lacI	b0345	170	1.34	Transcriptional repressor of the *lac* operon
→ lctR	b3604	14	4.76	Regulatory protein for L-Lactate dehydrogenase genes
nac	b1988	105	1.66	Nitrogen assimilation control protein
rnk	b0610	126	1.53	Regulator of nucleoside diphosphate kinase
rpoS	b2741	88	1.78	RNA polymerase sigma S factor
ttk	b3641	134	1.50	Putative transcriptional regulator
L : DNA replication. recombination and repair				
b0299	b0299	245	1.13	IS3 putative transposase
dinG	b0799	114	1.60	ATP-dependent helicase. SOS
dinI	b1061	61	2.02	Inhibits RecA-mediated self-cleavage. SOS
dinP	b0231	82	1.81	Putative tRNA synthetase. SOS
exo	b2798	78	1.84	5'-3' exonuclease. excision repair
intA	b2622	177	1.31	Prophage CP4-57 integrase
→ recA	b2699	51	2.30	DNA strand exchange and renaturation. SOS
recN	b2616	237	1.16	Recombination and DNA repair. SOS
sbmC	b2009	94	1.74	SbmC protein. SOS
xthA	b1749	207	1.23	Exonuclease III
METABOLISM				
C : Energy production and conversion				
aceA	b4015	195	1.25	Isocitrate lyase
aceB	b4014	120	1.56	Malate synthase A
aldA	b1415	67	1.93	Aldehyde dehydrogenase. NAD-linked
atpA	b3734	186	1.28	Membrane-bound ATP synthase alpha-subunit
cyoA	b0432	89	1.78	Cytochrome o ubiquinol oxidase subunit II

cyoC	b0430	26	3.59	Cytochrome o ubiquinol oxidase subunit III
cyoD	b0429	3	7.41	Cytochrome o ubiquinol oxidase subunit IV
dctA	b3528	74	1.89	Uptake of C4-dicarboxylic acids
fdhF	b4079	23	3.85	Subunit of formate dehydrogenase H.
fdoG	b3894	69	1.92	Formate dehydrogenase-O major subunit
glpD	b3426	178	1.31	Sn-glycerol-3-phosphate dehydrogenase
glpK	b3926	92	1.75	Glycerol kinase
glpQ	b2239	46	2.50	Glycerol-3-phosphate diesterase
→ mdh	b3236	53	2.19	Malate dehydrogenase
nifU	b2529	27	3.41	Formation/repair of [Fe-S] clusters present in iron-sulfur proteins
pckA	b3403	112	1.62	Phosphoenolpyruvate carboxykinase
sdhB	b0724	173	1.32	Succinate dehydrogenase. Iron sulfur protein
sdhD	b0722	131	1.52	Succinate dehydrogenase. Hydrophobic subunit
sixA	b2340	41	2.74	Phosphohistidine phosphatase affecting phosphorelay of ArcB
sucA	b0726	7	6.54	2-oxoglutarate dehydrogenase (decarboxylase)
sucB	b0727	83	1.81	2-oxoglutarate dehydrogenase (dihydrolipoyltranssuccinate)
sucD	b0729	43	2.69	Succinyl-CoA synthetase. Alpha subunit
xdhD	b2881	115	1.59	Putative dehydrogenase

G : Carbohydrate transport and metabolism

agp	b1002	85	1.80	Periplasmic glucose-1-phosphatase
gcd	b0124	197	1.25	Glucose dehydrogenase
glgS	b3049	168	1.35	Glycogen biosynthesis. rpoS dependent
glpX	b3925	140	1.47	Unknown function in glycerol metabolism
lamB	b4036	36	2.94	Maltose high-affinity receptor
malE	b4034	91	1.76	Periplasmic maltose-binding protein
malF	b4033	87	1.79	Part of maltose permease
malS	b3571	164	1.36	Alpha-amylase
mglA	b2149	215	1.21	ATP-binding galactose-binding transport protein
mglB	b2150	66	1.93	Galactose-binding transport protein
mrsA	b3176	111	1.62	Similar to phosphoglucomutases and phosphomannomutases
pgm	b0688	149	1.45	Phosphoglucomutase
→ rbsB	b3751	48	2.41	D-ribose periplasmic binding protein, chemotaxis
rbsC	b3750	159	1.41	D-ribose high-affinity transport system
rbsD	b3748	154	1.42	D-ribose high-affinity transport system
sfsA	b0146	64	1.97	Regulatory for maltose metabolism

E : Amino acid transport and metabolism

ansB	b2957	65	1.96	Periplasmic L-asparaginase II
argC	b3958	201	1.24	N-acetyl-gamma-glutamylphosphate reductase
argR	b3237	137	1.49	Repressor of arg regulon
gadA	b3517	30	3.15	Glutamate decarboxylase isozyme
idnD	b4267	175	1.31	L-idonate dehydrogenase
leuD	b0071	106	1.66	Isopropylmalate isomerase subunit
metH	b4019	72	1.90	Repressor of metE and metF
nifS	b2530	62	1.98	Cysteine desulfurase
putP	b1015	183	1.29	Major sodium/proline symporter

F : Nucleotide transport and metabolism : none

H : Coenzyme metabolism

metK	b2942	110	1.63	Methionine adenosyltransferase
pnuC	b0751	169	1.34	Required for NMN transport
ubiE	b3833	220	1.20	Ubiquinone/menaquinone biosynthesis methyltransferase

I : Lipid metabolism :

fabA	b0954	127	1.53	Trans-2-decenoyl-ACP isomerase
fadB	b3846	21	4.18	Fatty acid oxidation complex. 4-enzyme protein
fadE	b0221	84	1.80	Acyl-coenzyme A dehydrogenase
fadL	b2344	181	1.29	Transport of long-chain fatty acids
pgpA	b0418	102	1.67	Phosphatidylglycerophosphatase
pssA	b2585	239	1.15	Phosphatidylserine synthase. Phospholipid synthesis
uppS	b0174	156	1.41	Undecaprenyl pyrophosphate synthetase (peptidoglycan)

Q : Secondary metabolites biosynthesis. transport and metabolism

idnO	b4266	148	1.45	5-keto-D-gluconate 5-reductase
ucpA	b2426	50	2.32	Short-chain dehydrogenases/reductases (SDR) family

CELLULAR PROCESSES

D: Cell division and chromosomal partitioning

ftsL	b0083	17	4.34	Cell division and growth
sulA	b0958	33	3.07	Inhibits cell division and ftsZ ring formation. SOS

O : Post-translational modification. protein turnover. chaperones

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dnaJ	b0015	163	1.36	Chaperone with DnaK. Heat shock protein
dnaK	b0014	107	1.64	Chaperone Hsp70. Heat shock proteins
eco	b2209	9	6.17	Ecotin. Periplasmic serine protease inhibitor
fkpA	b3347	93	1.74	FKBP-type peptidyl-prolyl cis-trans isomerase (rotamase)
glnE	b3053	100	1.69	Adenylylating enzyme for glutamine synthetase
htpG	b0473	218	1.20	Chaperone Hsp90. Heat shock protein
htpX	b1829	76	1.88	Heat shock protein. Integral membrane protein
→ *msrA*	b4219	22	3.87	Peptide methionine sulfoxide reductase

M+N : Cell envelope biogenesis and secretion

amiB	b4169	243	1.13	N-acetylmuramoyl-l-alanine amidase II. Murein hydrolase
ddg	b2378	124	1.54	Putative heat shock protein
fhiA	b0229	71	1.91	Flagellar biosynthesis
→ *fimA*	b4314	28	3.29	Major type 1 subunit fimbrin (pilin)
fimI	b4315	49	2.33	Fimbrial protein
htrL	b3618	214	1.21	Involved in lipopolysaccharide biosynthesis
lepB	b2568	121	1.56	Leader peptidase (signal peptidase I)
mraW	b0082	238	1.15	Putative apolipoprotein
nlpB	b2477	191	1.26	Lipoprotein-34
nlpC	b1708	182	1.29	Lipoprotein
ompC	b2215	129	1.52	Outer membrane protein 1b
ompG	b1319	162	1.37	Outer membrane protein G
pspA	b1304	2	8.42	Phage shock protein. Inner membrane protein
pspB	b1305	59	2.04	Phage shock protein
pspC	b1306	12	5.58	Phage shock protein. Activates phage shock-protein expression
pspD	b1307	11	5.61	Phage shock protein
pspE	b1308	47	2.47	Phage shock protein
→ *pspF*	b1303	211	1.22	*psp* operon transcriptional activator
→ *tatE*	b0627	57	2.12	Membrane translocation of folded periplasmic proteins

P : Inorganic ion transport and metabolism

chaA	b1216	234	1.16	Sodium-calcium/proton antiporter
chaC	b1218	70	1.91	Accessory and regulatory protein for *chaA*
cutC	b1874	24	3.74	Copper homeostasis protein
cysP	b2425	74	1.89	Thiosulfate binding protein
cysU	b2424	136	1.49	Thiosulfate transport system permease
fur	b0683	188	1.27	Ferric iron uptake negative regulator
modA	b0763	99	1.70	Molybdate-binding periplasmic protein. Permease
modB	b0764	223	1.19	Molybdate transport permease protein
modC	b0765	216	1.20	ATP-binding component of molybdate transport
modE	b0761	139	1.47	Molybdate uptake regulatory protein
sodC	b1646	32	3.10	Superoxide dismutase precursor (Cu-Zn)
trkH	b3849	226	1.19	Potassium uptake

T : Signal transduction mechanism :

→ *cpxP*	b3914	1	22.9	Suppresses toxic envelope protein effects. CpxA/R activated
rpoE	b2573	63	1.98	Extra-cytoplasmic Sigma-E factor
rseA	b2572	55	2.15	Negative regulatory protein of sigma-E factor
rseB	b2571	29	3.20	Negative regulatory protein of sigma-E factor
→ *spy*	b1743	31	3.13	Periplasmic protein related to spheroblast formation

NOT CHARACTERIZED

R : Function unknown : General prediction only

→ *ycfJ*	b1110	5	6.95	Similarity to *Rickettsia* 17 kda surface antigen
→ *ycfR*	b1112	8	6.4	Exported / Outer membrane protein ?
yoaB	b1809	10	6.03	Putative translation initiation inhibitor
yebE	b1846	13	5.47	Similarity to an *Y.enterocolitica* protein
→ *yqcC*	b2792	15	4.45	Similarity to *E.carotovora* orf1 exoenzyme
yhhY	b3441	16	4.35	Putative acetyltransferase
→ *yggN*	b2958	21	4.1	Activated by RpoE
→ *yneA*	b1516	25	3.6	Putative periplasmic binding protein
ybeD	b0631	35	2.97	Homology to one histine kinase sensor domain of *M. grisea*
ydcI	b1422	37	2.83	Putative transcriptional regulator LysR-type
yddL	b1472	38	2.82	Putative outer membrane porin protein
→ *yccA*	b0970	39	2.76	Putative carrier/transport membrane protein. Degraded by FtsH
→ *yfcX*	b2341	40	2.75	Putative fatty oxidation complex alpha subunit
yjbO	b4050	44	2.58	Similarity to a putative exported *Y. pestis* protein
yrdD	b3283	45	2.54	Putative DNA topoisomerase
ybjF	b0859	56	2.14	Putative 23S rRNA (uracil-5-)-methyltransferase
yihN	b3874	60	2.03	Putative resistance protein (transport)
ycfT	b1115	68	1.93	Integral membrane protein

yeeF	b2014	77	1.84	Putative amino acid/amine transport protein
yfiE	b2577	86	1.79	Putative transcriptional regulator LysR-type
yeeD	b2012	90	1.77	Belongs to the UPF0033 family
yliH	b0836	95	1.73	Putative receptor
yfcM	b2326	96	1.73	Putative transporting ATPase
ybiX	b0804	97	1.73	Putative enzyme
yfhF/nifA	b2528	101	1.68	Putative regulator
ygfQ	b2884	113	1.62	Belongs to YicO/YieG/YjcD family
ybhR	b0792	118	1.58	Simlarity to E. coli YbhS, YhhJ and YhiG. IM protein
ybdH	b0599	130	1.52	Putative oxidoreductase
yihR	b3879	132	1.51	Putative aldose-1-epimerase
ydcT	b1441	138	1.47	Putative ATP-binding component of a transport system
ygiS	b3020	143	1.45	Putative transport periplasmic protein
ybaZ	b0454	150	1.44	Similarity to Cysteine methyltransferase
ydaM	b1341	155	1.41	Contains 1 GGDEF Dufl domain
tfaR	b1373	157	1.41	Phage lambda tail fiber gene homolog in prophage Rac
yceL	b1065	161	1.38	Belongs to the major fator family. Integral Membrane Protein
yheT	b3353	167	1.35	Belongs to the UPF0017 family
yjdC	b4135	172	1.33	Similarity to S. glaudescens TcmR
ybiW	b0823	174	1.32	Putative formate acetyltransferase
ybiF	b0813	176	1.31	Putative transmembrane subunit
ynaI	b1330	180	1.3	Belongs to the UPF0003 family. Integral membrane protein
yceE	b1053	185	1.28	Putative transport protein
yhdP	b1657	196	1.25	Putative transport protein
ygjE	b3063	213	1.21	Putative tartrate carrier
csiE	b2535	217	1.2	Stationary phase inducible protein
yfdE	b2371	219	1.2	Putative enzyme
yeeE	b2013	221	1.19	Putative transport system permease protein
yegQ	b2081	228	1.18	Putative peptidase (family U32)
glcA	b2975	229	1.17	Putative permease
yfdW	b2374	232	1.17	Putative enzyme
yfeT	b2427	233	1.17	Belongs to the Sis family, RpiR subfamily
ygjK	b3080	236	1.16	Putative isomerase
ydeW	b1512	242	1.13	Putative transcriptional regulator. SorC family

S : Function unknown

b1228	b1228	4	-7.04	Unknown
ycfL	b1104	6	6.70	Unknown
→yghO	b2981	18	4.31	Unknown
yiaH	b3561	19	4.18	Unknown. Integral membrane protein
→yceP	b1060	34	3.06	Unknown
yqeC	b2876	42	2.70	Unknown
□ygiB	b3037	54	2.15	Unknown. Integral membrane protein
ycfT	b1115	68	1.93	Unknown
yhjJ	b3527	73	1.90	Unknown
yceB	b1063	80	1.84	Unknown
ybiX	b0804	97	1.73	Unknown
ygiQ	b3015	98	1.71	Unknown
yagV	b0289	103	1.67	Unknown
yoeA	b1995	104	1.66	Unknown
ybhQ	b0791	108	1.64	Unknown
ybcI	b0527	109	1.63	Unknown
ybbF	b0524	116	1.59	Unknown
ybgI	b0710	117	1.58	Unknown
yncH	b1455	119	1.58	Unknown
yfbM	b0681	122	1.56	Unknown
yjiM	b4335	123	1.54	Unknown
yjfO	b4189	125	1.54	Unknown
ychN	b1219	128	1.53	Unknown
ynaC	b1373	141	1.47	Unknown
ymfE	b1138	142	1.46	Unknown
yfcN	b2331	144	1.45	Unknown
yrbC	b3192	145	1.45	Unknown
yfdQ	b2360	147	1.45	Unknown
yfeY	b2432	151	1.44	Unknown
ygiM	b3055	152	1.43	Unknown
yhgA	b3411	153	1.43	Unknown
yhjQ	b3534	158	1.41	Unknown

yfcF	b2301	160	1.39	Unknown
yfcI	b2305	165	1.35	Unknown
yjiD	b4326	171	1.34	Unknown
yfbP	b2275	179	1.30	Unknown
yphB	b2544	184	1.28	Unknown
yfbN	b2273	187	1.28	Unknown
ylbH	b0499	189	1.27	Unknown
ybhM	b0787	190	1.26	Unknown. Integral membrane protein
yrbL	b3207	192	1.26	Unknown
yjfY	b4199	193	1.25	Unknown
ynfA	b1582	194	1.25	Unknown
yajI	b0412	198	1.25	Unknown
yedI	b1958	200	1.24	Unknown
yafZ	b0252	202	1.24	Unknown
yjjU	b4377	203	1.24	Unknown
yfhH	b2561	204	1.24	Unknown
yafN	b0232	205	1.23	Unknown
yrbE	b3194	206	1.23	Unknown
yfgC	b2494	208	1.22	Unknown
yfjQ	b2633	209	1.22	Unknown
ycaK	b0901	210	1.22	Unknown
yfeS	b2420	212	1.22	Unknown
b4250	b4250	222	1.19	Unknown
ybgA	b0707	224	1.19	Unknown
yeeA	b2008	225	1.19	Unknown
ypfI	b2474	230	1.17	Unknown
b2394	b2394	231	1.17	IS186 hypothetical protein
yegK	b2072	235	1.16	Unknown
ybcJ	b0528	240	1.14	Unknown
yhiN	b3492	241	1.14	Unknown
ypfG	b2466	244	1.13	Unknown
ydiY	b1722	246	1.13	Unknown
yjjJ	b4385	247	1.12	Unknown
ycaP	b0906	248	1.11	Unknown
yfgJ	b2510	249	1.10	Unknown

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Table 4. Genes under-expressed in *E. coli* TG1 biofilm versus exponential growth phase.

The genes found to be under-expressed at a significant level (P-value ≤0.05) are indicated. They have been classified according to the COGs functional categories annotation system.

a : Gene names according to *E. coli* Colibri database.

b : Gene names according to Blattner nomenclature.

c : Rank position 1= the most repressed gene in *E. coli* biofilm.

d : Ratio of gene expression in *E. coli* biofilm versus gene expression in planktonic cultures.

e : Function description according to *E. coli* Colibri database.

Genes		Rank	Bio/Exp	Function - description
a	b	c	d	e
INFORMATION STORAGE AND PROCESSING				
J : Translation, ribosomal structure and metabolism				
def	b3287	62	0.67	Peptide deformylase
frr	b0172	81	0.70	Ribosome releasing factor
prfA	b1211	166	0.84	Peptide chain release factor RF-1
rbfA	b3167	40	0.61	Ribosome-binding factor A
rbn	b3886	176	0.87	tRNA processing exoribonuclease BN
rimJ	b1066	88	0.72	Acetylation of 30S ribosomal subunit protein S5
rpmG	b3636	79	0.69	50S ribosomal subunit protein L33
rpsV	b1480	161	0.83	30S ribosomal subunit protein S22
serS	b0893	96	0.73	Serine tRNA synthetase
thrS	b1719	63	0.67	Threonine tRNA synthetase
K : Transcription				
gcvR	b2479	69	0.67	Transcriptional regulation of *gcv* operon
malT	b3418	119	0.77	Positive regulator of *mal* regulon
osmE	b1739	66	0.67	Osmotically inducible lipoprotein E
oxyR	b3961	108	0.75	Activator of hydrogen peroxide-inducible genes
xylR	b3569	170	0.86	Putative regulator of *xyl* operon
L : DNA replication, recombination and repair				
dnaG	b3066	187	0.92	DNA primase
holA	b0640	134	0.79	DNA polymerase III delta subunit
hupB	b0440	41	0.61	DNA-binding protein HU-beta
intE	b1140	164	0.84	Prophage e14 integrase
nudG	b1759	174	0.87	CTP pyrophosphohydrolase
uvrD	b3813	183	0.91	DNA-dependent ATPase I and helicase II
xerC	b3811	109	0.76	Site-specific recombinase
METABOLISM				
C : Energy production and conversion				
adhE	b1241	1	0.23	Iron-dependent alcohol dehydrogenase
aldB	b3588	180	0.89	Aldehyde dehydrogenase
cydA	b0733	93	0.73	Cytochrome d terminal oxidase. Polypeptide subunit I
cydB	b0734	127	0.78	Cytochrome d terminal oxidase Polypeptide subunit II
dcuC	b0621	106	0.75	Transport of dicarboxylates
fumB	b4122	145	0.81	Fumarase B
icdA	b1136	136	0.80	Isocitrate dehydrogenase
pflB	b0903	29	0.56	Formate acetyltransferase 1
pta	b2297	120	0.77	Phosphotransacetylase
G : Carbohydrate transport and metabolism				
bglX	b2132	159	0.83	Beta-D-glucoside glucohydrolase
cmr	b0842	158	0.83	Proton motive force efflux pump

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cpsG	b2048	135	0.80	Phosphomannomutase
crr	b2417	70	0.68	Glucose-specific IIA component
eno	b2779	27	0.55	Enolase. Glycolysis
fba	b2925	48	0.63	Fructose-bisphosphate aldolase. Glycolysis
fbp	b4232	103	0.75	Fructose-bisphosphatase
fruA	b2167	61	0.66	Fructose-specific transport protein
fruB	b2169	71	0.68	Fructose-specific IIA/fpr component
fruK	b2168	33	0.57	Fructose-1-phosphate kinase
gapA	b1779	4	0.32	Glyceraldehyde-3-phosphate dehydrogenase A.
gpmA	b0755	36	0.60	Phosphoglyceromutase 1. Glycolysis
manY	b1818	18	0.47	PTS enzyme IIC. Mannose-specific
nagZ	b1107	94	0.73	Beta-hexosaminidase. Cell wall synthesis
pfkA	b3916	56	0.66	6-phosphofructokinase I. Glycolysis
pgk	b2926	53	0.65	Phosphoglycerate kinase. Glycolysis
ptsI	b2416	47	0.62	PEP-protein phosphotransferase system enzyme I
sgaH	b4196	90	0.72	Hexulose-6-phosphate synthase
sgaU	b4197	146	0.82	Hexulose-6-phosphate isomerase
shiA	b1981	144	0.81	Putative shikimate transport protein
torT	b0994	38	0.60	Part of regulation of *tor* operon.
tpiA	b3919	45	0.62	Triosephosphate isomerase. Glycolysis

E : Amino acid transport and metabolism

arcC	b0521	80	0.69	Putative carbamate kinase. Arginine degradation
argF	b0273	102	0.74	Ornithine carbamoyltransferase 2
aroG	b0754	11	0.43	DAHP synthetase. Aromatic amino acids biosynthesis
aroH	b1704	143	0.81	DAHP synthetase. Aromatic amino acids biosynthesis
asnB	b0674	148	0.82	Asparagine synthetase B
edd	b1851	162	0.84	6-phosphogluconate dehydratase
ggt	b3447	186	0.92	Gamma-glutamyltranspeptidase
glnA	b3870	58	0.66	Glutamine synthetase
glnB	b2553	20	0.49	Regulatory protein P-II for glutamine synthetase
hisB	b2022	25	0.53	Imidazole glycerolphosphate dehydratase
hisC	b2021	17	0.47	Histidinol-phosphate aminotransferase
hisG	b2019	165	0.84	ATP phosphoribosyltransferase
hisI	b2026	44	0.62	Phosphoribosyl-ATP pyrophosphatase
ilvL	b3766	14	0.45	ilvGEDA operon leader peptide
oppA	b1243	42	0.61	Oligopeptide transport. Periplasmic binding protein
oppB	b1244	74	0.68	Oligopeptide transport. Permease protein
oppC	b1245	129	0.78	Oligopeptide transport. Permease protein
oppD	b1246	65	0.67	ATP-binding protein of oligopeptide transport system
oppF	b1247	172	0.86	ATP-binding protein of oligopeptide transport system
pepQ	b3847	101	0.74	Proline dipeptidase
trpA	b1260	72	0.68	Tryptophan synthase. alpha protein
trpB	b1261	39	0.60	Tryptophan synthase. beta protein

F : Nucleotide transport and metabolism

cyaA	b3806	43	0.61	Adenylate cyclase
hpt	b0125	59	0.66	Purine salvage
tdk	b1238	113	0.77	Thymidine kinase

H : Coenzyme metabolism

bioH	b3412	118	0.77	Biotin biosynthesis
dxs	b0420	150	0.82	1-deoxyxylulose-5-phosphate synthase. Flavoprotein
folC	b2315	179	0.89	Dihydrofolate synthetase
mobA	b3857	49	0.63	Molybdopterin
tbpA	b0068	76	0.69	Thiamin-binding periplasmic protein

I : Lipid metabolism : none

Q : Secondary metabolites biosynthesis, transport and metabolism

pmbA	b4235	149	0.82	Maturation of antibiotic MccB17

CELLULAR PROCESSES

D: Cell division and chromosomal partitioning

zipA	b2412	126	0.78	Cell division protein involved in FtsZ ring

O : Post-translational modification, protein turnover, chaperones

clpA	b0882	133	0.79	ATP-binding component of serine protease
fkpB	b0028	50	0.64	Peptidyl-prolyl cis-trans isomerase (a rotamase)
ppiB	b0525	52	0.64	Peptidyl-prolyl cis-trans isomerase B (rotamase B)

M+N : Cell envelope biogenesis and secretion

cpsB	b2049	91	0.72	Colanic acid biosynthesis
dacA	b0632	34	0.58	D-alanyl-D-alanine carboxypeptidase

exbB	b3006	16	0.46	Uptake of enterochelin
exbD	b3005	15	0.46	Uptake of enterochelin
lpp	b1677	24	0.53	Murein lipoprotein
lpxD	b0179	84	0.71	Third step of endotoxin (lipidA) synthesis
pbpG	b2134	128	0.78	Penicillin-binding protein 7
sohB	b1272	60	0.66	Putative protease
yfbE	b2253	64	0.67	Putative enzyme

P : Inorganic ion transport and metabolism

bfd	b3337	10	0.41	Iron storage and mobility [2Fe-2S]
feoA	b3408	12	0.43	Ferrous iron transport protein A
feoB	b3409	57	0.66	ferrous iron transport protein B
fhuF	b4367	31	0.57	Ferric hydroxamate transport
focA	b0904	2	0.30	Formate transporter
hcaA1	b2538	86	0.72	Large subunit of phenylpropionate dioxygenase

T : Signal transduction mechanism : none

NOT CHARACTERIZED

R : Function unknown : General prediction only

yncE	b1452	5	0.34	Putative receptor
yhiX	b3516	8	0.39	Putative AraC-type regulatory protein
yfiD	b2579	13	0.44	Putative formate acetyltransferase
yodB	b1974	21	0.51	Putative cytochrome
ynfK	b1593	22	0.52	Putative dethiobiotin synthetase
ycgT	b1200	26	0.53	Putative dihydroxyacetone kinase
yebL	b1857	35	0.59	Putative high-affinity zinc uptake system protein
yeeX	b2007	37	0.60	Putative alpha helix protein
ykgM	b0296	51	0.64	Putative ribosomal protein
ybaO	b0447	54	0.65	Putative Lrp-like transcriptional regulator
etp	b0982	55	0.66	Putative protein-tyrosine-phosphatase
ygjH	b3074	67	0.67	Putative tRNA synthetase
yqhC	b3010	68	0.67	Putative AraC-type regulatory protein
yhcJ	b3223	73	0.68	Putative enzyme
yeiA	b2147	77	0.69	Putative oxidoreductase
ybgS	b0753	82	0.71	Putative homeobox protein
yhfW	b3380	85	0.71	Putative mutase
ydgF	b1600	98	0.74	Possible chaperone
ybcC	b0539	99	0.74	Putative exonuclease
ybjW	b0873	100	0.74	Putative prismane
yjiL	b4334	105	0.75	Putative enzyme
ctsA	b0598	112	0.76	Putative carbon starvation protein
ydjG	b1771	114	0.77	Hypothetical oxidoreductase
yeaU	b1800	116	0.77	Putative tartrate dehydrogenase
ygjU	b3089	117	0.77	Putative symporter protein
yejO	b2190	121	0.78	Putative ATP-binding component of a transport system
yeiC	b2166	124	0.78	Putative sugar kinase
ynjE	b1757	130	0.79	Putative thiosulfate sulfur transferase
yjbC	b4022	142	0.81	Putative pseudo-uridine synthase
yadF	*b0126*	147	0.82	Putative carbonic anhydrase
essD	*b0554*	151	0.82	Lysis protein homolog to lambdoid prophage DLP12
yneI	*b1525*	152	0.82	Putative aldehyde dehydrogenase
perM	*b2493*	156	0.83	Putative permease
yjjP	*b4364*	157	0.83	Putative structural protein
yhdX	*b3269*	160	0.83	Putative transport system permease protein
yihO	*b3876*	167	0.85	Putative permease
ycgS	*b1199*	169	0.85	Putative dihydroxyacetone kinase
yqhH	*b3014*	173	0.86	Putative lipoprotein
b0878	*b0878*	175	0.87	Putative membrane protein
ygfH	*b2920*	177	0.88	Putative coenzyme A transferase
yegH	*b2063*	178	0.88	Putative transport protein
ydiF	*b1694*	181	0.89	Putative enzyme
yeeZ	*b2016*	182	0.90	Putative enzyme of sugar metabolism
ydhM	*b1649*	185	0.91	Putative transcriptional regulator
ydjK	*b1775*	188	0.93	Putative transport protein

S : Function unknown

b3007	b3007	3	0.30	Unknown
yfjF	b2618	6	0.35	Unknown
ynaK	b1365	7	0.37	Unknown
b3004	b3004	9	0.39	Unknown

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yodA	b1973	19	0.48	Unknown
ymfA	b1122	23	0.52	Unknown
yjgD	b4255	28	0.56	Unknown
yeaQ	b1795	30	0.57	Unknown
yaiI	b0387	32	0.57	Unknown
yceD	b1088	46	0.62	Unknown
b0100	b0100	75	0.69	Unknown
ydcN	b1434	78	0.69	Unknown
ygiH	b3059	83	0.71	Unknown
ytfI	b4215	87	0.72	Unknown
ymfO	b1151	89	0.72	Unknown
ytfH	b4212	92	0.73	Unknown
ynhA	b1679	95	0.73	Unknown
ybaM	b0466	97	0.73	Unknown
ynfB	b1583	104	0.75	Unknown
ydgA	b1614	107	0.75	Unknown
yggJ	b2946	110	0.76	Unknown
yadS	b0157	111	0.76	Unknown
yfeK	b2419	115	0.77	Unknown
ycgR	b1194	122	0.78	Unknown
yfdS	b2362	123	0.78	Unknown
yadH	b0128	125	0.78	Unknown
yhhZ	b3442	131	0.79	Unknown
yhiJ	b3488	132	0.79	Unknown
ycbJ	b0919	137	0.81	Unknown
elaA	b2267	138	0.81	Unknown
ybhN	b1788	139	0.81	Unknown
ydgH	b1604	140	0.81	Unknown
yfjR	b2634	141	0.81	Unknown
ynfC	b1585	153	0.82	Unknown
yhgG	b3410	154	0.82	Unknown
ydjZ	b1752	155	0.83	Unknown
ydjY	b1751	163	0.84	Unknown
yhbV	b3159	168	0.85	Unknown
b2791	b2791	171	0.86	Unknown
ynjB	b1754	184	0.91	Unknown

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Table 5. Genes over-expressed (≥2) in *E. coli* TG1 biofilm versus both exponential and stationary growth phase.

a : Gene names according to *E. coli* Colibri database.

b : Gene names according to Blattner nomenclature.

c : Ratio of gene expression in *E. coli* biofilm versus exponential growth phase.

d : Ratio of gene expression in *E. coli* biofilm versus stationary growth phase.

e : Function description according to *E. coli* Colibri database.

Arrow: mutants where biofilm formation were reduced compared to wt.

The genes have been classified according to the COGs functional categories annotation system used by the NCBI.

Genes		Bio/Exp	Bio/Sta	Function - description
a	b	c	d	e
INFORMATION STORAGE AND PROCESSING				
J : Translation. ribosomal structure and metabolism				
rne	b1084	2.06	3.57	RNase E
K : Transcription				
→ lctR	b3604	4.76	8.07	Regulatory protein for L-Lactate dehydrogenase genes
L : DNA replication. recombination and repair				
dinI	b1061	2.02	2.92	Inhibits RecA-mediated self-cleavage. SOS
METABOLISM				
C : Energy production and conversion				
glpQ	b2239	2.50	2.15	Glycerol-3-phosphate diesterase
→ mdh	b3236	2.19	3.68	Malate dehydrogenase
sixA	b2340	2.74	3.24	Phosphohistidine phosphatase affecting phosphorelay of ArcB
G : Carbohydrate transport and metabolism				
lamB	b4036	2.94	3.73	Maltose high-affinity receptor
→ rbsB	b3751	2.41	3.83	D-ribose periplasmic binding protein. chemotaxis
E : Amino acid transport and metabolism				
gadA	b3517	3.15	4.84	Glutamate decarboxylase isozyme
F : Nucleotide transport and metabolism : none				
H : Coenzyme metabolism : none				
I : Lipid metabolism : none				
Q : Secondary metabolites biosynthesis. transport and metabolism : none				
CELLULAR PROCESSES				
D: Cell division and chromosomal partitioning : none				
O : Post-translational modification. protein turnover. Chaperones : none				
pspA	b1304	8.42	3.86	Phage shock protein. Inner membrane protein
pspB	b1305	2.04	3.44	Phage shock protein
pspC	b1306	5.58	2.55	Phage shock protein. Activates phage shock-protein expression
pspD	b1307	5.61	2.48	Phage shock protein
→ tatE	b0627	2.12	5.02	Membrane translocation of folded periplasmic proteins
P : Inorganic ion transport and metabolism : none				
T : Signal transduction mechanism :				
→ cpxP	b3914	22.9	13.15	Suppresses toxic envelope protein effects. CpxA/R activated
rseA	b2572	2.15	2.28	Negative regulatory protein of sigma-E factor
rpoE	b2573	1.98	5.86	Extracytoplasmic sigma E factor
→ spy	b1743	3.13	5.02	Periplasmic protein related to spheroblast formation

NOT CHARACTERIZED
R : Function unknown : General prediction only

yebE	b1846	5.47	3.14	Similarity to an *Y.enterocolitica* protein
→ yqcC	b2792	4.45	2.31	Similarity to *E.carotovora* orfl exoenzyme
→ yfcX	b2341	2.75	4.83	Putative fatty oxidation complex alpha subunit
yjbO	b4050	2.58	5.66	Similarity to a putative exported *Y. pestis* protein

S : Function unknown

→ yceP	b1060	3.06	6.46	Unknown
→ ygiB	b3037	2.15	2.09	Unknown

Table 6. Inactivation of the genes described in the study and TG1*gfp* strain construction: primers used in the linear DNA, 3 step PCR inactivation protocol.

5   [a] : Gene names according to *E. coli* Colibri database.

[b] : nomenclature according to Institute Pasteur database.

* : Genes inactived by a removable frt kanamycin cassette.

Target genes[a]	Primers name[b]	SEQ ID NO	Primers sequence
cpxP*	CpxP.A1.500-5	1	5'CGGCATCATTACGTCAAGCAAAAG3'
	CpxP.B1.500-3	2	5'GCGCCAGCGCCGCGAGGGACTCAG3'
	CpxP.B2.frtL-5	3	5'GAACTTCGGAATAGGAACTAATAGTAAACCCTGTTTTCCTTGCC3'
	CpxP.A2.frtL-3	4	5'*GAAGCAGCTCCAGCCTACACCATCATTTGCTCCCAAAATCTTTC3*'
	CpxP.ext-5		
	CpxP.ext-3	5	5'CCCGAATTCCGAAGTGCTTTTAATGTGTCG3'
		6	5'CGCCTGGATCTGTCATCGGTG3'
cpxR*	CpxR.A1.500-5	7	5'CGTGAGTTGCTACTACTCAATAG3'
	CpxR.B1.500-3	8	5'GCCGGACGAATCAGATAAAG3'
	CpxR.B2.frtL-5	9	5'GAACTTCGGAATAGGAACTAAGGTTTAAAACCTTGCGTGGTC3'
	CpxR.A2.frtL-3	10	5'GAAGCAGCTCCAGCCTACACGAAATTACGTCATCAGACGTCGC3'
	CpxR.ext-5	11	5'GATTGATTCATAAATACTCC3'
	CpxR.ext-3	12	5'CAAACAGTAAGTTAATGAAATC3'
cutC	*CutC.A1.500-5*	13	5'CACTATTGCATCAGAAGCGG3'
	CutC.B1.500-3	14	5'CCTTTCTGGTTCGAAAAGTGG3'
	CutC.B2.GBL-5	15	5'*CTTCACGAGGCAGACCTCAGCGCCTGATTTTTACCGTTGCATCATGTCGC3*'
	CutC.A2.GBL-3		
	CutC.ext-5	16	5'*GATTTTGAGACACAACGTGGCTTTCATTTTTACTCCTTAATTACGCCGAC3*'
	CutC.ext-3		
		17	5'GGAATACCTTACATTGATGA3'
		18	5'CTTTAGATGCCTTTAATTTAG3'
cyoC	CyoC.A1.500-5	19	5'CCATGCTGATGATTGCAGCC3'
	CyoC.B1.500-3	20	5'CCGACGCCACAACCAGTGAC3'
	CyoC.B2.GBL-5	21	5'CTTCACGAGGCAGACCTCAGCGCCTAATGAGTCATTCTACCGATCAC3'
	CyoC.A2.GBL-3	22	5'*GATTTTGAGACACAACGTGGCTTTCATTTTTCAGCCCTGCCTTAGTAATC3*'
	CyoC.ext-5		
	CyoC.ext-3	23	5'CAGGGATGACCTACTGGTGG3'
		24	5'GGATTCGCGCCAAACCACAG3'
dinI	DinI.A1.500-5	25	5'GTTTAACCGCAACCATATGC3'
	DinI.B1.500-3	26	5'CGATTCCTGCTTCTAATATC3'
	DinI.B2.GBL-5	27	5'CTTCACGAGGCAGACCTCAGCGCCTAATATGCAGTGATTTTTTTTGCC3'
	DinI.A2.GBL-3	28	5'*GATTTTGAGACACAACGTGGCTTTCATAATAGCCCCCTGTTGAA3*'
	DinI.ext-5		
	DinI.ext-3	29	5'CCTGACTGCGCTGAAAGTCG3'
		30	5'GACGCCGATACTCGTTTACC3'

ecO	EcO.A1.500-5	31	5'CGCCGCGTTGCAGAATGTTG3'
	EcO.B1.500-3	32	5'CCGGATGTGGCGTATGCTGATAAGACGC3'
	EcO.B2.GBL-5	33	5'CTTCACGAGGCAGACCTCAGCGCCCAACGCGGTAGTTCGCTAAACTGCCG3'
	EcO.A2.GBL-3	34	5'GATTTTGAGACACAACGTGGCTTTCCATTTTTTTTGCTTTCCTTC3'
	EcO.ext-5		
	EcO.ext-3	35	5'ATTTTTGAAATTAACGCTCG3'
		36	5'GTTGAAACCGCAACCCGTTC3'
fadB	FadB.A1.500-5	37	5'GATCACTTCCACATCTTCAG3'
	FadB.B1.500-3	38	5'GATTTCATTTTTAAATGCGG3'
	FadB.B2.GBL-5	39	5'CTTCACGAGGCAGACCTCAGCGCCTAAGGAGTCACAATGGAACAGGTTG3'
	FadB.A2.GBL-3	40	5'GATTTTGAGACACAACGTGGCTTTCATGTCAGTCTCCTGAATCC3'
	FadB.ext-5		
	FadB.ext-3	41	5'CTGGCCTCAATACCCAGTTG3'
		42	5'GTTTACTGGATCAAACGCCGGACGC3'
fdhF	FdhF.A1.500-5	43	5'GTCTGCAAACGCTCAACGAC3'
	FdhF.B1.500-3	44	5'GTCGTTCTCCAGATCTTCCG3'
	FdhF.B2.GBL-5	45	5'CTTCACGAGGCAGACCTCAGCGCCTAATACCGTCCTTTCTACAG3'
	FdhF.A2.GBL-3	46	5'GATTTTGAGACACAACGTGGCTTTCCATCGGTCTCGCTCCAGTTAATC3'
	FdhF.ext-5		
	FdhF.ext-3	47	5'GCCGCTGTTTGACGGTGGAC3'
		48	5'CGCCCAGTACTCGGAATAAC3'
gadA	GadA.A1.500-5	49	5'CCTTTGAACCGTTGGGGCTG3'
	GadA.B1.500-3	50	5'CTTATCTACTCGAATTTGGC3'
	GadA.B2.GBL-5	51	5'CTTCACGAGGCAGACCTCAGCGCCGATAACATAACGTTGTAAAAAC3'
	GadA.A2.GBL-3	52	5'GATTTTGAGACACAACGTGGCTTTCATTTCGAACTCCTTAAATTTATTTG3'
	GadA.ext-5		
	GadA.ext-3	53	5'GTTGCGCGGAGATGAAAATG3'
		54	5'CATGAAGATTTAATGCCTCC3'
lctR	LctR.A1.500-5	55	5'GCACTGCTCTCGATTGTCTG3'
	LctR.B1.500-3	56	5'GGGCCGCTCATACCTGAATG3'
	LctR.B2.GBL-5	57	5'CTTCACGAGGCAGACCTCAGCGCCTGATTATTTCCGCAGCCAGCGAT3'
	LctR.A2.GBL-3	58	5'GATTTTGAGACACAACGTGGCTTTCCATTAAGGAATCATCCACGTTAAG3'
	LctR.ext-5		
	LctR.ext-3	59	5'GGTGGCGCGCTGTATGAGTG3'
		60	5'CCTAAATCATGTGGACGACC3'
malM to *malG*	MalMG.A1.500-5	61	5'ACGACTCCAGCGGATCGCGCGGCAAC3'
	MalMG.B1.500-3	62	5'CAATAGTGGAATTGTTGCTTTATC3'
	MalMG.B2.GBL-5	63	5'CTTCACGAGGCAGACCTCAGCGCCTAGCCCTTGTGGAGGTTCCTGCAAT3'
	MalMG.A2.GBL-3	64	5'GATTTTGAGACACAACGTGGCTTTCATTTCTCATCCTTGTTTTATC3'
	MalMG.ext-5		
	MalMG.ext-3	65	5'GGTTTTCGACCAGTTTGACTAAG3'
		66	5'CGTTGGTGCTGTTAGCACTGTATC3'
mdh	Mdh.A1.500-5	67	5'GCATAAGTCACCCGATATGGTGG3'
	Mdh.B1.500-3	68	5'CTCGCTGGGCGAACTGATGGG3'
	Mdh.B2.GBL-5	69	5'CTTCACGAGGCAGACCTCAGCGCCTAATTGATTAGCGGATAATAAAAAAC3'
	Mdh.A2.GBL-3	70	5'GATTTTGAGACACAACGTGGCTTTCATCCTAAACTCCTTATTATATTG3'
	Mdh.ext-5		
	Mdh.ext-3	71	5'CTGCAACGCGGCGACGATTTC3'
		72	5'GGCAAAACTTCCTCCAAACCG3'
nifS	NifS.A1.500-5	73	5'CCTTTCTTATCTGGAACAAC3'
	NifS.B1.500-3	74	5'CGCCCAGACGCAGGCCAAAC3'
	NifS.B2.GBL-5	75	5'CTTCACGAGGCAGACCTCAGCGCCTAATCGGTATCGGAATCAG3'
	NifS.A2.GBL-3	76	5'GATTTTGAGACACAACGTGGCTTTCATTGCTCTATAAACTCCGTACATCAC3'
	NifS.ext-5		
	NifS.ext-3	77	5'CATGAGACTGACATCTAAAG3'
		78	5'CTTCTTTTACGAAGTCCAGC3'

nifU	NifU.A1.500-5	79	5'CATCGCAAAAGAAGAGATGG3'
	NifU.B1.500-3	80	5'CTCAGCGCCTGGGTATCGAG3'
	NifU.B2.GBL-5	81	5'CTTCACGAGGCAGACCTCAGCGCCTAAGAGTTGAGGTTTGGTTATG3'
	NifU.A2.GBL-3	82	5'*GATTTTGAGACACAACGTGGCTTTCA*TTATAAATTCTCCTGATTC3'
	NifU.ext-5		
	NifU.ext-3	83	5'GGTGCGCTGTATGTACGTCG3'
		84	5'GGTTAATGGTTGCAGATTGC3'
nlpE	NlpE.A1.500-5	85	5'ACATGTTGCTATTCCCGATG3'
	NlpE.B1.500-3	86	5'GCAGTGTGGGCGAAGGAGAC3'
	NlpE.B2.GBL-5	87	5'CACGAGGCAGACCTCAGCGCTAACCCGTCTTGAGACAGAAACAAAC3'
	NlpE.A2.GBL-3	88	5'*TTGAGACACAACGTGGCTTTCA*TCCATTCCTTCTTTTTATTCCCG3'
	NlpE.ext-5		
	NlpE.ext-3	89	5'ATCTTTCCGTCTGGTATCTG3'
		90	5'GACTCGCCAGATGTGCTCAC3'
pspA to pspE	PspAE.A1.500-5	91	5'CCCGAGCTCACCATCATCGGTGCCGTAGCGAG3'
	PspAE.B1.500-3	92	5'GATAATCAATTACCGAAAAGCCATC3'
	PspAE.B2.GBL-5	93	5'CTTCACGAGGCAGACCTCAGCGCCTAAAAGAATTCACCATGAGCGG3'
	PspAE.A2.GBL-3	94	5'*GATTTTGAGACACAACGTGGCTTTCC*ATAATGTTGTCCTCTTGATTTCTG3'
	PspAE.ext-5		
	PspAE.ext-3	95	5'CAGTTCACCGTACTCAATCACGC3'
		96	5'CGAGTTGCTGAATATCCTGCCACTCC3'
rbsB	RbsB.A1.500-5	97	5'GGTATTGGTCGTCCGCTGGG3'
	RbsB.B1.500-3	98	5'CGCTCACGTTGCGCTTCCAC3'
	RbsB.B2.GBL-5	99	5'CTTCACGAGGCAGACCTCAGCGCCTAGTTTTAATCAGGTTGTATG3'
	RbsB.A2.GBL-3	100	5'*GATTTTGAGACACAACGTGGCTTTCA*TATTCAAGATGTCCTGTAG3'
	RbsB.ext-5		
	RbsB.ext-3	101	5'GGCGTGACCATGGTTTATAC3'
		102	5'GAAGTTCGCGAGCCGGAGCC3'
rpoE	RpoE.A1.500-5	103	5'GACCTGATGCTGGTCAGCCAGGCGTAG3'
	RpoE.B1.500-3	104	5'CGCTTCAGAAGGTACTCCCAG3'
	RpoE.B2.GBL-5	105	5'CTTCACGAGGCAGACCTCAGCGCCCAGGCGTTGACGATAGCGGG3'
	RpoE.A2.GBL-3	106	5'*GATTTTGAGACACAACGTGGCTTTCA*TCCGAGGTAAAGTCTCCCCA3'
	RpoE.ext-5		
	RpoE.ext-3	107	5'GAACCTTCCGTTACCGGGCCTTTAC3'
		108	5'GCAACATTGCATTAATGCGACGAC3'
rseA *	RseA.A1.500-5	109	5'GCATAAAGTGGCGAGTCTGG3'
	RseA.B1.500-3	110	5'GTAATTTCGATTCGGTGTCC3'
	RseA.B2.frtL-5	111	5'GAACTTCGGAATAGGAACTAAGTTTGAGCAGGCGCAAACCCAGC3'
	RseA.A2.frtL-3	112	5'*GAAGCAGCTCCAGCC*TACACCATGCCTAATACCCTTATCC3'
	RseA.ext-5		
	RseA.ext-3	113	5'GGTCCTGGTTGAACGGGTCC3'
		114	5'GTTCCAGCGTTTCACCATCG3'
rseB	RseB.A1.500-5	115	5'CCATTTCGATATCTCTTCAC3'
	RseB.B1.500-3	116	5'CGTCCTCGCATTTGTTATGC3'
	RseB.B2.GBL-5	117	5'CTTCACGAGGCAGACCTCAGCGCCATGATCAAAGAGTGGGCTAC3'
	RseB.A2.GBL-3	118	5'*GATTTTGAGACACAACGTGGCTTTCA*TTACTGCGATTGCGTTCC3'
	RseB.ext-5		
	RseB.ext-3	119	5'CTTAATCCGTGACTCAATGC3'
		120	5'GAAATGTTCATACCGTATGG3'
sixA	SixA.A1.500-5	121	5'CGCACCGCAGGTTGCTGAAC3'
	SixA.B1.500-3	122	5'GTGATGTTTTCACTCCCCTGATTC3'
	SixA.B2.GBL-5	123	5'CTTCACGAGGCAGACCTCAGCGCCTGATGAGTTCCAAATTATGC3'
	SixA.A2.GBL-3	124	5'*GATTTTGAGACACAACGTGGCTTTCA*TATTGCACCGCTTTTGTTAACCAG3'
	SixA.ext-5		
	SixA.ext-3	125	5'GCTGATTGGCACACAAGGGC3'
		126	5'CATTGATTCAGTCAATAGCCAATG3'

sodC	SodC.A1.500-5	127	5'GCAATCACGTCTGCCGTTTACC3'
	SodC.B1.500-3	128	5'GATCGGATGCTCGTAAAAGCC3'
	SodC.B2.GBL-5	129	5'CTTCACGAGGCAGACCTCAGCGCCCCGATCAACCTAAACCGCTGGG3'
	SodC.A2.GBL-3	130	5'GATTTTGAGACACAACGTGGCTTTCATAGGACCTCCGTTCATTG3'
	SodC.ext-5		
	SodC.ext-3	131	5'CGTTCAAACATCTGCATCAGAG3'
		132	5'GGCGTCGCGTTGGCGTGGTTAG3'
spy	Spy.A1.500-5	133	5'GACACGCTGAATTTTATGCC3'
	Spy.B1.500-3	134	5'CTGCCCTGCCGTCAGTTTCG3'
	Spy.B2.GBL-5	135	5'CTTCACGAGGCAGACCTCAGCGCCTAATCTTTCAGCCAAAAAACTTAAGAC3'
	Spy.A2.GBL-3	136	5'GATTTTGAGACACAACGTGGCTTTCCATATTCTATATCCTTCCTTTC3'
	Spy.ext-5		
	Spy.ext-3	137	5'GTCGGTATCGTGAGAACACC3'
		138	5'CTTACAGACATCCAGGCGTG3'
sucA	SucA.A1.500-5	139	5'GGCTTGTTAGCGGCATATCG3'
	SucA.B1.500-3	140	5'GACACGTTTTTCACTACGTG3'
	SucA.B2.GBL-5	141	5'CTTCACGAGGCAGACCTCAGCGCCTAAATAAAGGATACACAATG3'
	SucA.A2.GBL-3	142	5'GATTTTGAGACACAACGTGGCTTTCATCGTGATCCCTTAAGCATC3'
	SucA.ext-5		
	SucA.ext-3	143	5'CGCGAGCATTTACAGATGCC3'
		144	5'GCTTCACCGTACTGCTTACG3'
sulA	SulA.A1.500-5	145	5'CAGCTTCAGTTGATTTCGCC3'
	SulA.B1.500-3	146	5'CAGTTGGTTTTCATGGGTCG3'
	SulA.B2.GBL-5	147	5' CTTCACGAGGCAGACCTCAGCGCCTAAGTAAATTTAGGATTAATCCTG3'
	SulA.A2.GBL-3	148	5'GATTTTGAGACACAACGTGGCTTTCCATAATCAATCCAGCCCCTG3'
	SulA.ext-5		
	SulA.ext-3	149	5' GCAAATCTTTCAGTCTTTCC3'
		150	5' CATTTCAAAGCCAACATACG3'
tatE	TatE.A1.500-5	151	5'GTCTGATGACCTGTTATGAC3'
	TatE.B1.500-3	152	5'CAACGCCACCAGATGTGTTC3'
	TatE.B2.GBL-5	153	5'CTTCACGAGGCAGACCTCAGCGCCTGACGTGGCGAGCAGGACGC3'
	TatE.A2.GBL-3	154	5'GATTTTGAGACACAACGTGGCTTTCATAGATACCTTCTTGAC3'
	TatE.ext-5		
	TatE.ext-3	155	5'TGATGCTGGTAATGAAATCG3'
		156	5'CGCGGTCGTATGGATCGTGC3'
ybeD	YbeD.A1.500-5	157	5'TACTTTTAAAGGCCGTGAAG3'
	YbeD.B1.500-3	158	5'GCCCGAGGATGCGCTTCTAT3'
	YbeD.B2.GBL-5	159	5'CTTCACGAGGCAGACCTCAGCGCCTAACTCGCTTCTCCGTTAC3'
	YbeD.A2.GBL-3	160	5'GATTTTGAGACACAACGTGGCTTTCATGTCAGCTCCGGCGTAAC3'
	YbeD.ext-5		
	YbeD.ext-3	161	5'CGGACACACTGACAAAGCAG3'
		162	5'CCATATTGACGTTTAATGCC3'
ybjF	YbjF.A1.500-5	163	5'TCATGGAAGACGAAACGTTG3'
	YbjF.B1.500-3	164	5'CGGAAGTGAAAACTGTCTCT3'
	YbjF.B2.GBL-5	165	5'CTTCACGAGGCAGACCTCAGCGCCTAAAAAAGCCGCATGTG3'
	YbjF.A2.GBL-3	166	5'GATTTTGAGACACAACGTGGCTTTCATACATTGACCTTCACATC3'
	YbjF.ext-5		
	YbjF.ext-3	167	5'CAACCTGGCTACATAATGCC3'
		168	5'GATACCTACAAAACGTTTGC3'
yccA	YccA.A1.500-5	169	5'CGGGCGGTGGGGATGTTTAG3'
	YccA.B1.500-3	170	5'CAGTGGTTAAAGAGTGGCGG3'
	YccA.B2.GBL-5	171	5'CTTCACGAGGCAGACCTCAGCGCCTAATCTCACCCGCTAACAC3'
	YccA.A2.GBL-3	172	5'GATTTTGAGACACAACGTGGCTTTCATTGAGTCACTCTCTATG3'
	YccA.ext-5		
	YccA.ext-3	173	5'CTGCACTGGCGCACGTCGCC3'
		174	5'CGATGGCAGCGTGGAAGTGG3'

yceP	YceP.A1.500-5	175	5'GCGAAAACTTCTCCATTGCC3'
	YceP.B1.500-3	176	5'CAGCGGGCCATAATCCCTTG3'
	YceP.B2.GBL-5	177	5'CTTCACGAGGCAGACCTCAGCGCCTAACATGACATGACCATCC3'
	YceP.A2.GBL-3	178	5'GATTTTGAGACACAACGTGGCTTTCATCATGGCCCCCTAATTCG3'
	YceP.ext-5		
	YceP.ext-3	179	5'CCAGTATATTCAACAGGGGG3'
		180	5'CTTCGCCAGTTGGATCCAGG3'
ycfJ	YcfJ.A1.500-5	181	5'CAGGCTGCACACCAGATGGC3'
	YcfJ.B1.500-3	182	5'CGGAATTTACCAACAAAGAG3'
	YcfJ.B2.GBL-5	183	5'CTTCACGAGGCAGACCTCAGCGCCTAACAAGGCTGTACTCTG3'
	YcfJ.A2.GBL-3	184	5'GATTTTGAGACACAACGTGGCTTTCACGGGAACACCTCCTTC3'
	YcfJ.ext-5		
	YcfJ.ext-3	185	5'CAGACATTTACGCTATTGGC3'
		186	5'GGACCTCGTCGAAGCGACCG3'
ycfL	YcfL.A1.500-5	187	5'GATATATACGGCAGCAAAAC3'
	YcfL.B1.500-3	188	5'GGCAATGCCTATGGCTTTAC3'
	YcfL.B2.GBL-5	189	5'CTTCACGAGGCAGACCTCAGCGCCTAAGGGGTGAATCTTGATG3'
	YcfL.A2.GBL-3	190	5'GATTTTGAGACACAACGTGGCTTTCATCGTTACAGACCTTTATG3'
	YcfL.ext-5		
	YcfL.ext-3	191	5'GCGATTATATTTAGTGTGCG3'
		192	5'CTGACCAGATAATTTCGCCC3'
ycfR	YcfR.A1.500-5	193	5'CAGCTGTGCTTCATGCTTAG3'
	YcfR.B1.500-3	194	5'GCCGGCTGGACTGGATAACC3'
	YcfR.B2.GBL-5	195	5'CTTCACGAGGCAGACCTCAGCGCCTAAGCATTAACCCTCATT3'
	YcfR.A2.GBL-3	196	5'GATTTTGAGACACAACGTGGCTTTCATAATAGTGGCCTTATGC3'
	YcfR.ext-5		
	YcfR.ext-3	197	5'CATGAAGCAGCCTGCCGGGG3'
		198	5'GACAAACGTGCAAACCCAAC3'
ydcI	YdcI.A1.500-5	199	5'GTCGAATGTACCGGCACCCC3'
	YdcI.B1.500-3	200	5'CATCAACAGTATTGCTTTCC3'
	YdcI.B2.GBL-5	201	5'CTTCACGAGGCAGACCTCAGCGCCTGAAAGGTGAAGGGATCTGTC3'
	YdcI.A2.GBL-3	202	5'GATTTTGAGACACAACGTGGCTTTCATAAGCGATGTTAAAAAC3'
	YdcI.ext-5		
	YdcI.ext-3	203	5'GCGTGTCGTATTCTTCTTGC3'
		204	5'CGCTTCATCTCACTGAGGAC3'
yebE	YebE.A1.500-5	205	5'CAAAAAATTGTCGGTCAGGC3'
	YebE.B1.500-3	206	5'GCATATTCACAGCCTGGTTC3'
	YebE.B2.GBL-5	207	5'CTTCACGAGGCAGACCTCAGCGCCTAATTCCGCTCTCTGGATAG3'
	YebE.A2.GBL-3	208	5'GATTTTGAGACACAACGTGGCTTTCATATTTGCTCCTCAATAAC3'
	YebE.ext-5		
	YebE.ext-3	209	5'GTGAAGATCTGGATGCTGCC3'
		210	5'GGTGTTATCGGGCGTAATCG3'
yfcX	YfcX.A1.500-5	211	5'CGCAAACACGGAACGGTAAC3'
	YfcX.B1.500-3	212	5'GAGATCACCAGTACCGAAGC3'
	YfcX.B2.GBL-5	213	5'CTTCACGAGGCAGACCTCAGCGCCTAAGAAGGTCAAAGCTATATGAA3'
	YfcX.A2.GBL-3	214	5'GATTTTGAGACACAACGTGGCTTTCATTATTCCGCCTCCAGAACCA3'
	YfcX.ext-5		
	YfcX.ext-3	215	5'GGTGATGACTGCCTTTATCC3'
		216	5'CATCTTCAGATTACACGGGC3'
yggN	YggN.A1.500-5	217	5'GAACCGTAGCCGTCGTCTGC3'
	YggN.B1.500-3	218	5'CATCGTGTCGGTACCGTGGG3'
	YggN.B2.GBL-5	219	5'CTTCACGAGGCAGACCTCAGCGCCTAATCCTCTATTTTAAGACG3'
	YggN.A2.GBL-3	220	5'GATTTTGAGACACAACGTGGCTTTCATAGTCTTCCCTCAAG3'
	YggN.ext-5		
	YggN.ext-3	221	5'GTGATGTCTTCTATTGACGG3'
		222	5'GTTGGCGGAGGCTTTATCAG3'

yghO	YghO.A1.500-5	223	5'CGACCAAGGTGCCTTGAGTC3'
	YghO.B1.500-3	224	5'GCAGCCGCGAACGCTGTACG3'
	YghO.B2.GBL-5	225	5'CTTCACGAGGCAGACCTCAGCGCCTAATACCAGCTAACTCAGGTTC3'
	YghO.A2.GBL-3	226	5'*GATTTTGAGACACAACGTGGCTTT*ATTAAGGAAGGTGCGAACAAGTC3'
	YghO.ext-5		
	YghO.ext-3	227	5'CTGCTCTTTGTTCTTGGTCG3'
		228	5'GCGCAGGGTCGCGATTCTCG3'
ygiB	YgiB.A1.500-5	229	5'GCGATGGAAGCGGGCTACTC3'
	YgiB.B1.500-3	230	5'GTTCACGCAGCTCAACGAAG3'
	YgiB.B2.GBL-5	231	5'CTTCACGAGGCAGACCTCAGCGCCTGATACCGATGGAAAGAGTC3'
	YgiB.A2.GBL-3	232	5'*GATTTTGAGACACAACGTGGCTTT*CATTTTTGTCTTCCGGGACC3'
	YgiB.ext-5	233	5'GAATGGTTAACTCGCAGGTG3'
	YgiB.ext-3	234	5'CCTGATCCTGTAAATCCGTG3'
yhhY	YhhY.A1.500-5	235	5'CGCTGGTGAAATGGATATGG3'
	YhhY.B1.500-3	236	5'GATAAAAAAGCGCCTCTTAG3'
	YhhY.B2.GBL-5	237	5'CTTCACGAGGCAGACCTCAGCGCCTAAGATAGTGCCCTTTTTCTG3'
	YhhY.A2.GBL-3	238	5'*GATTTTGAGACACAACGTGGCTTT*CATTCCTTTGTCCTCTTTGG3'
	YhhY.ext-5		
	YhhY.ext-3	239	5'GTTTCGCGTACTCGAAATGG3'
		240	5'CGATAAGATGTTGACAGAGG3'
yiaH	YiaH.A1.500-5	241	5'GGAAAAAGCAGGGCTTAACG3'
	YiaH.B1.500-3	242	5'GTCAAATGCGTTTGTTTCGC3'
	YiaH.B2.GBL-5	243	5'CTTCACGAGGCAGACCTCAGCGCCTAAGTAAAAGCCCGGTCACATTGGAC3'
	YiaH.A2.GBL-3	244	5'*GATTTTGAGACACAACGTGGCTTT*CATCTGTGTCTCTGTATCTG3'
	YiaH.ext-5		
	YiaH.ext-3	245	5'CAAGCCCTGGAAGGTCCTGG3'
		246	5'CATATCTGCCAGTTAGTTGC3'
yjbO	YjbO.A1.500-5	247	5'CGATTAACGGTGGTATCAAG3'
	YjbO.B1.500-3	248	5'CCGTGGGCAGAGACACCTGG3'
	YjbO.B2.GBL-5	249	5'CTTCACGAGGCAGACCTCAGCGCCTAAGGGATTGTGCGGATGATCACAAC3'
	YjbO.A2.GBL-3	250	5'*GATTTTGAGACACAACGTGGCTTT*CATGATGCTCTCCCAAATATG3'
	YjbO.ext-5		
	YjbO.ext-3	251	5'GCAAAGGCGAGTGTGAGATG3'
		252	5'GAGCGGTTAAAAGAGATCAC3'
yneA	YneA.A1.500-5	253	5'GGCTGCATAAAACCCATGCC3'
	YneA.B1.500-3	254	5'CGACTGATGTTCATATTCGC3'
	YneA.B2.GBL-5	255	5'CTTCACGAGGCAGACCTCAGCGCCTGATGTGCATTACTTAACCG3'
	YneA.A2.GBL-3	256	5'*GATTTTGAGACACAACGTGGCTTT*CATGAAGATATCCTTTATGG3'
	YneA.ext-5		
	YneA.ext-3	257	5'GCTAACCTGGATGTGCTGGG3'
		258	5'GGTACCGGACATCCGGCAAC3'
yoaB	YoaB.A1.500-5	259	5'CCGGCAGATCGCCCCCCGCC3'
	YoaB.B1.500-3	260	5'GGTGTTGGCGCTGATACATC3'
	YoaB.B2.GBL-5	261	5'CTTCACGAGGCAGACCTCAGCGCCTAAGCTTTATCGAAGCAAAATAAG3'
	YoaB.A2.GBL-3	262	5'*GATTTTGAGACACAACGTGGCTTT*CATCATTTTGTCCTCATTATAC3'
	YoaB.ext-5		
	YoaB.ext-3	263	5'CCACGCCTGTGAATCTTCCG3'
		264	5'CCAGGGTTCCAGCCTTCCTG3'
yqcC	YqcC.A1.500-5	265	5'CTGTAAGCGCCTTGTAAGAC3'
	YqcC.B1.500-3	266	5'CGAAGCTGATGTTTGCGTCC3'
	YqcC.B2.GBL-5	267	5'CTTCACGAGGCAGACCTCAGCGCCTAATGCTGGAAATACTCTATC3'
	YqcC.A2.GBL-3	268	5'*GATTTTGAGACACAACGTGGCTTT*CATAAAGCAACCTCAATAAG3'
	YqcC.ext-5		
	YqcC.ext-3	269	5'CTTAAGCCTCTTCTGTAATC3'
		270	5'GGCCCGCGTGAATAGTCAGC3'

yqeC	YqeC.A1.500-5	271	5'GGGGATGCCATTATGGAGTG3'
	YqeC.B1.500-3	272	5'CACCAAACGACTCAGCATGG3'
	YqeC.B2.GBL-5	273	5'CTTCACGAGGCAGACCTCAGCGCCTAGCGGCCCGGGTATTCCGGG3'
	YqeC.A2.GBL-3	274	5'GATTTTGAGACACAACGTGGCTTTCACGAGTCTTTATGACCTC3'
	YqeC.ext-5	275	5'CTGCATTTTCTATTTCGACG3'
	YqeC.ext-3	276	5'GAACCTTGCGACGACTTGCC3'
λatt-*gfp*	ATT.A1.500-5	277	5'CGATGGCGATAATATTTCACC3'
	ATT.B1.500-3	278	5'CCCTGATACTCACCAGGCATC3'
	ATT.B2.xfp-5	279	5'TGAGTAGGACAAATCCGCCGCTAAAAAAGCAGGCTTCAAC3'
	ATT.A2.xfp-3	280	5'GCGTTTTTTATTGGTGAGAATTACTAACTTGAGCGAAACG3'
	ATT-ext5	281	5'GGCGATAAATTGCCGCATCG3'
	ATT-ext3	282	5'TGCCACCATCAAGGGAAAGCCC3'

Table 7. Primers used for the Q-RT-PCR experiments.

Primers were designed to amplify about 200-bp internal gene sequence.

Target genes	Primers name	SEQ ID NO:	Primers sequence
cpxP	cpxP-RT-5	283	5' CGCTGGCAGTCAGTTCATTAAGCC 3'
	cpxP-RT-3	284	5' GTCTCCAGTTCGCTAACATTAAC 3'
cyoD	cyoD-RT-5	285	5' CTACCGATCACAGCGGCGCGTCCC 3'
	cyoD-RT-3	286	5' GTTCCAGCCTTCATCTGATTTGG 3'
fimA	fimA-RT-5	287	5' CTGGCAATCGTTGTTCTGTCGGCTC 3'
	fimA-RT-3	288	5' GCTCCTTCCTGTGCCAGCGATGCG 3'
sucA	sucA-RT-5	289	5' GAACAGCTCTATGAAGACTTCTTAAC 3'
	sucA-RT-3	290	5' GCTGCAGGACTTTAACCTGCTTCACAT 3'
ycfJ	ycfJ-RT-5	291	5' GTTGGCGGGTATCGGGATTGGTGTC 3'
	ycfJ-RT-3	292	5' GTAATGCGATTTTCATCCTGCACC 3'
ycfR	ycfR-RT-5	293	5' CCCTCATCGCTGCGGCGATTTTAAGC 3'
	ycfR-RT-3	294	5' CCGGTTACAGAAGTAATACGGAAAG 3'
yebE	yebE-RT-5	295	5' GGCTGCTGGTCGCAAATAAATCAG 3'
	yebE-RT-3	296	5' GCAAGGATCAAACGTGCTGTACGC 3'